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International application number: PCT/US05/012337

International filing date: 12 April 2005 (12.04.2005)

Document type: Certified copy of priority document

Document details: Country/Office: US
Number: 60/563,285
Filing date: 16 April 2004 (16.04.2004)

Date of receipt at the International Bureau: 20 May 2005 (20.05.2005)

Remark: Priority document submitted or transmitted to the International Bureau in compliance with Rule 17.1(a) or (b)



World Intellectual Property Organization (WIPO) - Geneva, Switzerland
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APPLICATION NUMBER: 60/563,285

FILING DATE: *April 16, 2004*

RELATED PCT APPLICATION NUMBER: *PCT/US05/12337*



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PROVISIONAL APPLICATION FOR PATENT COVER SHEET

This is a request for filing a PROVISIONAL APPLICATION FOR PATENT under 37 CFR 1.53(c).

INVENTOR(S)				
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<input type="checkbox"/> Additional inventors are being named on the _____ separately numbered sheets attached hereto				
TITLE OF THE INVENTION (280 characters max)				
CANCER TREATMENT METHOD				
Direct all correspondence to: CORRESPONDENCE ADDRESS				
<input checked="" type="checkbox"/> Customer Number <input type="text" value="23347"/>		<div>Place Customer Number Bar Code Label here</div>		
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ENCLOSED APPLICATION PARTS (check all that apply)				
<input checked="" type="checkbox"/> Specification	Number of Pages	<input type="text" value="69"/>	<input type="checkbox"/> CD(s), Number	<input type="text"/>
<input checked="" type="checkbox"/> Drawing(s)	Number of Sheets	<input type="text" value="4"/>	<input type="checkbox"/> Other (specify)	<input type="text"/>
<input type="checkbox"/> Application Data Sheet. See 37 CFR 1.76				
METHOD OF PAYMENT OF FILING FEES FOR THIS PROVISIONAL APPLICATION FOR PATENT (check one)				
<input type="checkbox"/>	A check or money order is enclosed to cover the filing fees			FILING FEE AMOUNT (\$)
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<input checked="" type="checkbox"/> No.				
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Respectfully submitted,

SIGNATURE

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Date

11e Apr 04

REGISTRATION NO.

52,218

(if appropriate)

Docket Number:

PR60682P

USE ONLY FOR FILING A PROVISIONAL APPLICATION FOR PATENT

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PR60682Serial No.
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Invention: **CANCER TREATMENT METHOD**

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CANCER TREATMENT METHOD

BACKGROUND OF THE INVENTION

5 The present invention relates to pharmaceutical combinations and methods of treating cancer utilizing the same. Specifically, the invention relates to a combination of an inhibitor of VEGFR, and an inhibitor of Erb-B2 and/or Erb-B1, as well as use of the combination in the treatment of cancer.

10 Effective treatment of hyperproliferative disorders, including cancer, is a continuing goal in the oncology field. Protein tyrosine kinases catalyse the phosphorylation of cell growth and differentiation of specific tyrosyl residues in various proteins involved in the regulation of cell growth and differentiation. (A.F. Wilks, Progress in Growth Factor Research, 1990, 2, 97-111; S.A. Courtneidge, Dev. Suppl., 1993, 57-64; J.A. Cooper, Semin. Cell Biol., 1994, 5(6), 377-387; R.F. Paulson, Semin. Immunol., 1995, 7(4), 267-277; A.C. Chan, Curr. Opin. Immunol., 1996, 8(3), 394-401). Inappropriate or uncontrolled activation of many of such kinases, i.e., aberrant protein tyrosine kinase activity, for example by over-expression or mutation, has been shown to result in uncontrolled cell growth.

20 In cancer the growth of solid tumors has been shown to be dependent on angiogenesis. The progression of leukemias as well as the accumulation of fluid associated with malignant ascites and pleural effusions also involve pro-angiogenic factors. (See Folkmann, J., J. Nat'l. Cancer Inst., 1990, 82, 4-6.) Consequently, the targeting of pro-angiogenic pathways is a strategy being widely pursued in order to provide new therapeutics in these areas of great, unmet medical need.

30 Central to the process of angiogenesis are vascular endothelial growth factor (VEGF) and its receptors, termed vascular endothelial growth factor receptor(s) (VEGFRs). Three PTK receptors for VEGF have been identified: VEGFR1 (Flt-1); VEGFR2 (Flk-1 and KDR) and VEGFR3 (Flt-4). These receptors are involved in angiogenesis and participate in signal transduction. (Mustonen, T. et al J. Cell Biol. 1995:129:895-898; Ferrara and Davis-Smyth, Endocrine Reviews, 18(1):4-25, 1997; McMahon, G., The Oncologist, Vol. 5, No 90001, 3-10, April 2000).

Of particular interest is VEGFR2, which is a transmembrane receptor PTK expressed primarily in endothelial cells. Activation of VEGFR-2 by VEGF is a critical step in the signal transduction pathway that initiates tumor angiogenesis.

5 Consequently, antagonism of the VEGFR2 kinase domain would block phosphorylation of tyrosine residues and serve to disrupt initiation of angiogenesis, thereby providing a potent treatment for cancer or other disorders associated with inappropriate angiogenesis.

10 The erbB family of protein tyrosine kinases is one another group of such kinases which has been implicated in human malignancies. Elevated EGFR Erb B1 receptor activity has, for example, been implicated in non-small cell lung, bladder, renal cell, and head and neck cancers. Increased c-erbB-2 activity is associated with breast, ovarian, gastric and pancreatic cancers. Consequently, inhibition of such
15 protein tyrosine kinases should provide a treatment for disorders characterized by aberrant Erb family protein kinase activity.

International Patent Application PCT/US01/49367 filed December 19, 2001, and published as WO 02/059110 on August 1, 2002, discusses PTKs including
20 VEGFR family PTKs. This published application discloses bicyclic heteroaromatic compounds, including 5-({4-[(2,3-dimethyl-2*H*-indazol-6-yl)(methyl)amino]pyrimidin-2-yl}amino)-2-methylbenzenesulfonamide; as well as hydrochloride salts thereof. These compounds show inhibition activity against VEGFR-2. International Patent Application PCT/EP99/00048 filed January 8, 1999, and published as WO 99/35146
25 on July 15, 1999, discusses PTKs including erbB family PTKs. This published application discloses bicyclic heteroaromatic compounds, including N-{3-Chloro-4-[(3-fluorobenzyl)oxy]phenyl}-6-[5-({2-(methanesulphonyl)ethyl} amino)methyl]-2-furyl]-4-quinazolinamine; (4-(3-Fluoro-benzyloxy)-3-chlorophenyl) -(6-(2-((2-methanesulphonyl-ethylamino)-methyl)-thiazol-4-yl) quinazolin-4-yl)-amine; (4-(3-
30 Fluoro-benzyloxy)-3-bromophenyl)-(6-(5-((2-methanesulphonyl-ethyl amino)-methyl)-furan-2-yl)quinazolin-4-yl)-amine as well as hydrochloride salts thereof. These compounds show inhibition activity against erbB family PTKs.

Combination therapy is rapidly becoming the norm in cancer treatment, rather
35 than the exception. Oncologists are continually looking for anti-neoplastic compounds which when utilized in combination provides a more effective and/or

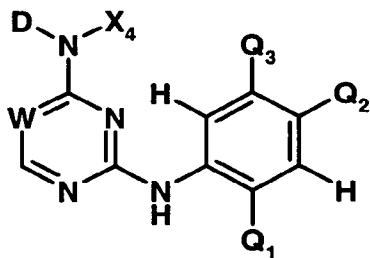
enhanced treatment to the individual suffering the effects of cancer. Typically, successful combination therapy provides improved and even synergistic effect over monotherapy.

Combination of VEGF and ErbB inhibitors have been explored in several pre-clinical tumor models (Ciardiello F. et al. Clin Cancer Res 2000; 6(9):3739-3747; Baker CH et al. Cancer Res 2002; 62(7):1996-2003; Shaheen RM et al. Br J Cancer 2001; 85(4):584-589; Jung YD et al. Eur J Cancer 2002; 38(8):1133-1140). In mice bearing human colon carcinoma xenografts, combined treatment with anti-EGFR mAb (c225) and VEGF antisense significantly improved survival compared to either agent alone (Ciardiello F. et al. Clin Cancer Res 2000; 6(9):3739-3747). Similarly, combination of antibodies against ErbB1 and VEGF receptor resulted in decreased angiogenesis and ascites formation compared to either antibody alone in a mouse model of peritoneal carcinomatosis (Baker CH et al. Cancer Res 2002; 62(7):1996-2003).

The present inventors have now identified combinations of chemotherapeutic agents that provide increased activity over monotherapy. In particular, multiple drug combinations that include inhibitors of the VEGFR family of kinases in combination with inhibitors of the ErbB family of kinases are described.

SUMMARY OF THE INVENTION

Briefly, one aspect of the present invention provides a method of treating cancer in a mammal, comprising: administering to said mammal a compound of formula I,

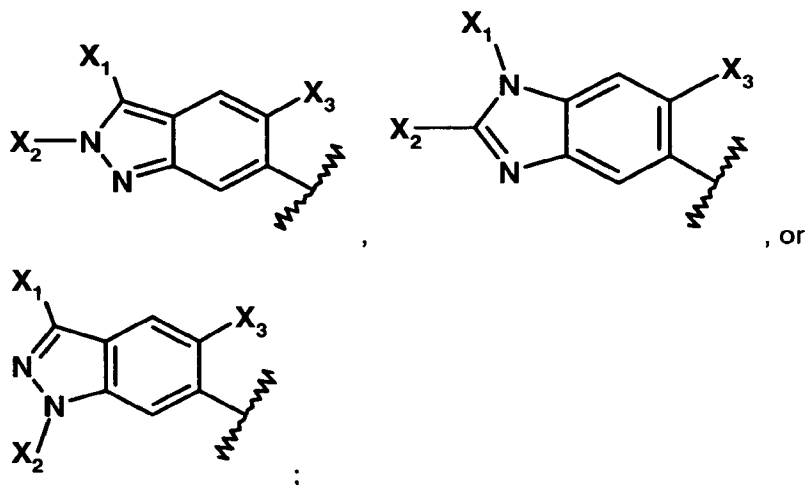


(I)

or a salt, solvate, or physiologically functional derivative thereof:

wherein:

D is



X_1 is hydrogen, C_1 - C_4 alkyl, C_1 - C_4 haloalkyl, or C_1 - C_4 hydroxyalkyl;

X_2 is hydrogen, C_1 - C_4 alkyl, C_1 - C_4 haloalkyl, $C(O)R^1$, or aralkyl;

X_3 is hydrogen or halogen;

10 X_4 is hydrogen, C_1 - C_4 alkyl, C_1 - C_4 haloalkyl, heteroaralkyl, cyanoalkyl, $-(CH_2)_pC=CH(CH_2)_tH$, $-(CH_2)_pC\equiv C(CH_2)_tH$, or C_3 - C_7 cycloalkyl;

p is 1, 2, or 3;

t is 0 or 1;

W is N or C-R, wherein R is hydrogen, halogen, or cyano;

15 Q_1 is hydrogen, halogen, C_1 - C_2 haloalkyl, C_1 - C_2 alkyl, C_1 - C_2 alkoxy, or C_1 - C_2 haloalkoxy;

Q_2 is A^1 or A^2 ;

Q_3 is A^1 when Q_2 is A^2 and Q_3 is A^2 when Q_2 is A^1 ;

wherein

20 A^1 is hydrogen, halogen, C_1 - C_3 alkyl, C_1 - C_3 haloalkyl, $-OR^1$, and

A^2 is the group defined by $-(Z)_m-(Z^1)-(Z^2)$, wherein

Z is CH_2 and m is 0, 1, 2, or 3, or

Z is NR^2 and m is 0 or 1, or

Z is oxygen and m is 0 or 1, or

25 Z is CH_2NR^2 and m is 0 or 1;

Z^1 is $S(O)_2$, $S(O)$, or $C(O)$; and

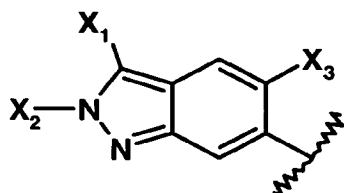
Z² is C₁-C₄ alkyl, NR³R⁴, aryl, arylamino, aralkyl, aralkoxy, or heteroaryl,

R¹ is C₁-C₄ alkyl;

R², R³, and R⁴ are each independently selected from hydrogen, C₁-C₄ alkyl, C₃-C₇ cycloalkyl, -S(O)₂R⁵, and -C(O)R⁵;

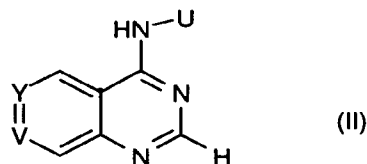
R⁵ is C₁.C₄ alkyl, or C₃.C₇ cycloalkyl; and

when Z is oxygen then Z^1 is $S(O)_2$ and when D is



then X₂ is C₁-C₄ alkyl, C₁-C₄ haloalkyl, C(O)R¹, or aralkyl; and

10 a compound of formula II



or a salt, solvate, or physiologically functional derivative thereof;

wherein

15 Y is CR^1 and V is N;

or Y is CR¹ and V is CR²;

R¹ represents a group CH₃SO₂CH₂CH₂NHCH₂-Ar-, wherein Ar is selected from phenyl, furan, thiophene, pyrrole and thiazole, each of which may optionally be

20 substituted by one or two halo, C₁₋₄ alkyl or C₁₋₄ alkoxy groups;

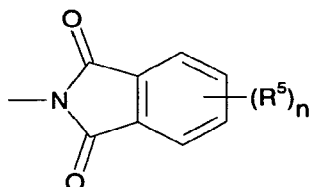
R² is selected from the group comprising hydrogen, halo, hydroxy, C₁₋₄ alkyl, C₁₋₄ alkoxy, C₁₋₄ alkylamino and di[C₁₋₄ alkyl]amino;

25 U represents a phenyl, pyridyl, 3H-imidazolyl, indolyl, isoindolyl, indolinyl, isoindolinyl, 1H-indazolyl, 2,3-dihydro-1H-indazolyl, 1H-benzimidazolyl, 2,3-dihydro-1H-benzimidazolyl or 1H-benzotriazolyl group, substituted by an R³ group and optionally substituted by at least one independently selected R⁴ group;

R^3 is selected from a group comprising benzyl, halo-, dihalo- and trihalobenzyl, benzoyl, pyridylmethyl, pyridylmethoxy, phenoxy, benzyloxy, halo-, dihalo- and trihalobenzyloxy and benzenesulphonyl;

5 or R^3 represents trihalomethylbenzyl or trihalomethylbenzyloxy;

or R^3 represents a group of formula



10 wherein each R^5 is independently selected from halogen, C_{1-4} alkyl and C_{1-4} alkoxy; and n is 0 to 3; and

each R^4 is independently hydroxy, halogen, C_{1-4} alkyl, C_{2-4} alkenyl, C_{2-4} alkynyl, C_{1-4} alkoxy, amino, C_{1-4} alkylamino, di[C_{1-4} alkyl]amino, C_{1-4} alkylthio, C_{1-4} alkylsulphinyl, C_{1-4} alkylsulphonyl, C_{1-4} alkylcarbonyl, carboxy, carbamoyl, C_{1-4} alkoxy carbonyl, C_{1-4} alkanoylamino, N-(C_{1-4} alkyl)carbamoyl, N,N-di(C_{1-4} alkyl)carbamoyl, cyano, nitro and trifluoromethyl.

20 In another aspect, the present invention provides a pharmaceutical composition comprising a compound of formula I or salt, solvate, or physiologically functional derivative thereof, and a compound of formula II or salt, solvate, or physiologically functional derivative thereof, optionally in association with a pharmaceutically acceptable diluent or carrier.

25 In a further aspect, the present invention provides a pharmaceutical combination comprising a compound of formula I or salt, solvate or physiologically functional derivative thereof and a compound of formula II or salt, solvate or physiologically functional derivative thereof for use in therapy.

30 In a further aspect, the present invention provides a pharmaceutical combination comprising a compound of formula I or salt, solvate or physiologically functional derivative thereof and a compound of formula II or salt, solvate or

physiologically functional derivative thereof for the manufacture of a medicament for the treatment of cancer.

5

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 depicts anti-tumor activity in a subcutaneous human xenograft mouse model dosed with a compound of Example 1 and a compound of Example 2 individually and in combination versus BT474 (human breast tumor line) from a representative experiment.

Table 1 depicts anti-tumor activity in a subcutaneous human xenograft mouse model dosed with a compound of Example 1 and a compound of Example 2 individually and in combination versus BT474 (human breast tumor line) in two independent experiments.

Figure 2 depicts anti-tumor activity in a subcutaneous human xenograft mouse model dosed with a compound of Example 1 and a compound of Example 2 individually and in combination versus NCI-H322 (non-small cell lung carcinoma).

Table 2 depicts anti-tumor activity in a subcutaneous human xenograft mouse model dosed with a compound of Example 1 and a compound of Example 2 individually and in combination versus NCI-H322 (non-small cell lung carcinoma) in three independent experiments.

DETAILED DESCRIPTION OF THE INVENTION

30

As used herein the term "neoplasm" refers to an abnormal growth of cells or tissue and is understood to include benign, i.e., non-cancerous growths, and malignant, i.e., cancerous growths. The term "neoplastic" means of or related to a neoplasm.

35

As used herein the term "agent" is understood to mean a substance that produces a desired effect in a tissue, system, animal, mammal, human, or other subject. Accordingly, the term "anti-neoplastic agent" is understood to mean a substance producing an anti-neoplastic effect in a tissue, system, animal, mammal, human, or other subject. It is also to be understood that an "agent" may be a single compound or a combination or composition of two or more compounds.

As used herein, the term "effective amount" means that amount of a drug or pharmaceutical agent that will elicit the biological or medical response of a tissue, system, animal or human that is being sought, for instance, by a researcher or clinician. Furthermore, the term "therapeutically effective amount" means any amount which, as compared to a corresponding subject who has not received such amount, results in improved treatment, healing, prevention, or amelioration of a disease, disorder, or side effect, or a decrease in the rate of advancement of a disease or disorder. The term also includes within its scope amounts effective to enhance normal physiological function.

As used herein, the term "lower" refers to a group having between one and six carbons.

As used herein, the term "alkyl" refers to a straight or branched chain hydrocarbon having from one to twelve carbon atoms, optionally substituted with substituents selected from the group consisting of lower alkyl, lower alkoxy, lower alkylsulfanyl, lower alkylsulfenyl, lower alkylsulfonyl, oxo, mercapto, amino optionally substituted by alkyl, carboxy, carbamoyl optionally substituted by alkyl, aminosulfonyl optionally substituted by alkyl, nitro, or lower perfluoroalkyl, multiple degrees of substitution being allowed. Examples of "alkyl" as used herein include, but are not limited to, n-butyl, n-pentyl, isobutyl, and isopropyl, and the like.

As used herein, the term "alkylene" refers to a straight or branched chain divalent hydrocarbon radical having from one to ten carbon atoms, optionally substituted with substituents selected from the group which includes lower alkyl, lower alkoxy, lower alkylsulfanyl, lower alkylsulfenyl, lower alkylsulfonyl, oxo, hydroxy, mercapto, amino optionally substituted by alkyl, carboxy, carbamoyl optionally substituted by alkyl, aminosulfonyl optionally substituted by alkyl, nitro, cyano, halogen and lower perfluoroalkyl, multiple degrees of substitution being

allowed. Examples of "alkylene" as used herein include, but are not limited to, methylene, ethylene, n-propylene, n-butylene, and the like.

As used herein, the term "C_{x-y}" where x and y represent an integer value refer to the number of carbon atoms in a particular chemical term to which it is attached. For instance, the term "C₁₋₄ alkyl" refers to an alkyl group, as defined herein, containing at least 1, and at most 4 carbon atoms. The term "C₁₋₄ alkylene" refers to an alkylene group, as defined above, which contains at least 1, and at most 4 carbon atoms.

10

As used herein, the term "C₁₋₄ haloalkyl" refers to a straight or branched chain hydrocarbon containing at least 1, and at most 4, carbon atoms substituted with at least one halogen. Examples of branched or straight chained "C₁₋₄ haloalkyl" groups useful in the present invention include, but are not limited to, methyl, ethyl, propyl, isopropyl, isobutyl and n-butyl substituted independently with one or more halogens, *e.g.*, fluoro, chloro, bromo and iodo.

15

As used herein, the term "C₁₋₄ hydroxyalkyl" refers to a straight or branched chain hydrocarbon containing at least 1, and at most 4, carbon atoms substituted with at least one hydroxy. Examples of branched or straight chained "C₁₋₄ hydroxyalkyl" groups useful in the present invention include, but are not limited to, methyl, ethyl, propyl, isopropyl, isobutyl and n-butyl substituted independently with one or more hydroxy groups.

20

As used herein, the term "C₃₋₇ cycloalkyl" refers to a non-aromatic cyclic hydrocarbon ring having from three to seven carbon atoms, which optionally includes a C₁₋₄ alkylene linker through which it may be attached. Exemplary "C₃₋₇ cycloalkyl" groups include, but are not limited to, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl and cycloheptyl.

25

30

As used herein, the term "heterocyclic" or the term "heterocyclyl" refers to a three to twelve-membered non-aromatic ring being unsaturated or having one or more degrees of unsaturation containing one or more heteroatomic substitutions selected from S, SO, SO₂, O, or N, optionally substituted with substituents selected from the group consisting of lower alkyl, lower alkoxy, lower alkylsulfanyl, lower alkylsulfenyl, lower alkylsulfonyl, oxo, hydroxy, mercapto, amino optionally

35

substituted by alkyl, carboxy, carbamoyl optionally substituted by alkyl, aminosulfonyl optionally substituted by alkyl, nitro, cyano, halogen, or lower perfluoroalkyl, multiple degrees of substitution being allowed. Such a ring may be optionally fused to one or more of another "heterocyclic" ring(s) or cycloalkyl ring(s). Examples of "heterocyclic" include, but are not limited to, tetrahydrofuran, pyran, 1,4-dioxane, 1,3-dioxane, piperidine, pyrrolidine, morpholine, tetrahydrothiopyran, tetrahydrothiophene, and the like.

As used herein, the term "aryl" refers to an optionally substituted benzene ring or to an optionally substituted benzene ring system fused to one or more optionally substituted benzene rings to form, for example, anthracene, phenanthrene, or naphthalene ring systems. Exemplary optional substituents include lower alkyl, lower alkoxy, lower alkylsulfonyl, lower alkylsulfenyl, lower alkylsulfonyl, oxo, hydroxy, mercapto, amino optionally substituted by alkyl, carboxy, tetrazolyl, carbamoyl optionally substituted by alkyl, aminosulfonyl optionally substituted by alkyl, acyl, aroyl, heteroaroyl, acyloxy, aroyloxy, heteroaroyloxy, alkoxycarbonyl, nitro, cyano, halogen, lower perfluoroalkyl, heteroaryl, or aryl, multiple degrees of substitution being allowed. Examples of "aryl" groups include, but are not limited to, phenyl, 2-naphthyl, 1-naphthyl, biphenyl, as well as substituted derivatives thereof.

As used herein, the term "aralkyl" refers to an aryl or heteroaryl group, as defined herein including both unsubstituted and substituted versions thereof, attached through a lower alkylene linker, wherein lower alkylene is as defined herein. As used herein, the term "heteroaralkyl" is included within the scope of the term "aralkyl". The term heteroaralkyl is defined as a heteroaryl group, as defined herein, attached through a lower alkylene linker, lower alkylene is as defined herein. Examples of "aralkyl", including "heteroaralkyl", include, but are not limited to, benzyl, phenylpropyl, 2-pyridinylmethyl, 4-pyridinylmethyl, 3-isoxazolylmethyl, 5-methyl-3-isoxazolylmethyl, and 2-imidazolyl ethyl.

As used herein, the term "arylamino" refers to an aryl or heteroaryl group, as defined herein, attached through an amino group $-NR^2$ -, wherein R^2 is as defined herein.

As used herein, the term "heteroaryl" refers to a monocyclic five to seven membered aromatic ring, or to a fused bicyclic aromatic ring system comprising two

of such monocyclic five to seven membered aromatic rings. These heteroaryl rings contain one or more nitrogen, sulfur, and/or oxygen heteroatoms, where N-oxides and sulfur oxides and dioxides are permissible heteroatom substitutions and may be optionally substituted with up to three members selected from a group consisting of lower alkyl, lower alkoxy, lower alkylsulfanyl, lower alkylsulfenyl, lower alkylsulfonyl, oxo, hydroxy, mercapto, amino optionally substituted by alkyl, carboxy, tetrazolyl, carbamoyl optionally substituted by alkyl, aminosulfonyl optionally substituted by alkyl, acyl, aroyl, heteroaroyl, acyloxy, aroyloxy, heteroaroyloxy, alkoxycarbonyl, nitro, cyano, halogen, lower perfluoroalkyl, heteroaryl, or aryl, multiple degrees of substitution being allowed. Examples of "heteroaryl" groups used herein include furan, thiophene, pyrrole, imidazole, pyrazole, triazole, tetrazole, thiazole, oxazole, isoxazole, oxadiazole, thiadiazole, isothiazole, pyridine, pyridazine, pyrazine, pyrimidine, quinoline, isoquinoline, benzofuran, benzothiophene, indole, indazole, and substituted versions thereof.

15

As used herein, the term "alkoxy" refers to the group R_aO- , where R_a is alkyl as defined above and the term " C_{1-2} alkoxy" refers to the group R_aO- , where R_a is C_{1-2} alkyl as defined above.

20

As used herein, the term "haloalkoxy" refers to the group R_aO- , where R_a is haloalkyl as defined above and the term " C_{1-2} haloalkoxy" refers to the group R_aO- , where R_a is C_{1-2} haloalkyl as defined above.

As used herein the term "aralkoxy" refers to the group R_bR_aO- , where R_a is alkylene and R_b is aryl, both as defined above.

25

As used herein the term "cyanoalkyl" refers to the group $-R_aCN$ wherein R_a is C_{1-3} alkylene as defined above. Exemplary "cyanoalkyl" groups useful in the present invention include, but are not limited to, cyanomethyl, cyanoethyl, and cyanopropyl.

30

As used herein, the term "aminosulfonyl" refers to the group $-SO_2NH_2$.

As used herein, the term "aroyl" refers to the group $R_aC(O)-$, where R_a is aryl as defined herein.

35

As used herein, the term "heteroaroyl" refers to the group $R_aC(O)-$, where R_a is heteroaryl as defined herein.

As used herein, the term "alkoxycarbonyl" refers to the group $R_aOC(O)-$,
5 where R_a is alkyl as defined herein.

As used herein, the term "acyloxy" refers to the group $R_aC(O)O-$, where R_a is alkyl, cycloalkyl, or heterocyclyl as defined herein.

10 As used herein, the term "aroyloxy" refers to the group $R_aC(O)O-$, where R_a is aryl as defined herein.

As used herein, the term "heteroaroyloxy" refers to the group $R_aC(O)O-$,
where R_a is heteroaryl as defined herein.

15

As used herein, the term "physiologically functional derivative" refers to any pharmaceutically acceptable derivative of a compound of formulae I or II, for example, an ester or an amide, which upon administration to a mammal is capable of providing (directly or indirectly) a compound of formulae I or II or an active metabolite
20 thereof. Such derivatives are clear to those skilled in the art, without undue experimentation, and with reference to the teaching of Burger's Medicinal Chemistry And Drug Discovery, 5th Edition, Vol 1: Principles and Practice, which is incorporated herein by reference to the extent that it teaches physiologically functional derivatives.

25 As used herein, the term "solvate" refers to a complex of variable stoichiometry formed by a solute (in this invention, a compound of formula I or II or a salt or physiologically functional derivative thereof) and a solvent. Such solvents for the purpose of the invention may not interfere with the biological activity of the solute. Examples of suitable solvents include, but are not limited to, water, methanol,
30 ethanol and acetic acid. Preferably the solvent used is a pharmaceutically acceptable solvent. Examples of suitable pharmaceutically acceptable solvents include water, ethanol and acetic acid. Most preferably the solvent used is water.

It is also noted that the compounds of formula I or II may form tautomers. It
35 is understood that all tautomers and mixtures of tautomers of the compounds of the

present invention, more specifically, the compounds of formula I or II are included within the scope of the compounds of the present invention.

5 The compounds of formulae I and II have the ability to crystallize in more than one form, a characteristic, which is known as polymorphism, and it is understood that such polymorphic forms ("polymorphs") are within the scope of formulae I and II. Polymorphism generally can occur as a response to changes in temperature or pressure or both and can also result from variations in the crystallization process. Polymorphs can be distinguished by various physical
10 characteristics known in the art such as x-ray diffraction patterns, solubility, and melting point.

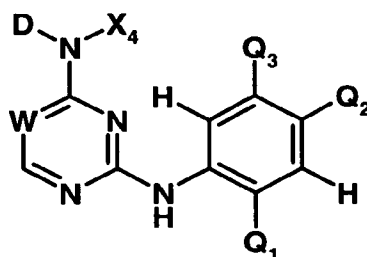
Typically, the salts of the compounds of formulae I and II are pharmaceutically acceptable salts. Salts encompassed within the term
15 "pharmaceutically acceptable salts" refer to non-toxic salts of the compounds of this invention. Salts of the compounds of formulae I and II may comprise acid addition salts derived from a nitrogen on a substituent in the compounds of formulae I and II. Representative salts include the following salts: acetate, benzenesulfonate, benzoate, bicarbonate, bisulfate, bitartrate, borate, bromide, calcium edetate, camsylate, carbonate, chloride, clavulanate, citrate, dihydrochloride, edetate,
20 edisylate, estolate, esylate, fumarate, gluceptate, gluconate, glutamate, glycolylarsanilate, hexylresorcinate, hydrabamine, hydrobromide, hydrochloride, hydroxynaphthoate, iodide, isethionate, lactate, lactobionate, laurate, malate, maleate, mandelate, mesylate, methylbromide, methylnitrate, methylsulfate, monopotassium maleate, mucate, napsylate, nitrate, N-methylglucamine, oxalate,
25 pamoate (embonate), palmitate, pantothenate, phosphate/diphosphate, polygalacturonate, potassium, salicylate, sodium, stearate, subacetate, succinate, tannate, tartrate, teoclate, tosylate and ditosylate, triethiodide, trimethylammonium and valerate. Other salts, which are not pharmaceutically acceptable, may be useful
30 in the preparation of compounds of this invention and these form a further aspect of the invention. Furthermore, such salt may be in anhydrous or hydrated form.

In one embodiment, the compound of formula I is a hydrochloride salt, preferably a monohydrochloric salt.

In one embodiment, the compound of formula II is a hydrochloride or ditosylate salt, preferably a ditosylate salt, more preferably the monohydrate of the ditosylate salt.

- 5 As recited above, a method of treating cancer is provided which includes administering a compound of formula I or a salt, solvate or physiologically functional derivative thereof and a compound of formula II or a salt, solvate or physiologically functional derivative thereof.

- 10 In one embodiment of the present invention is a compound of the formula I:



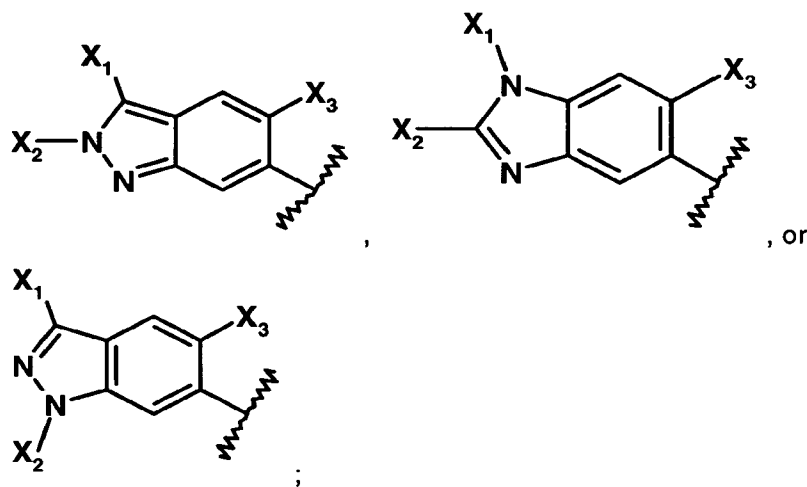
(I)

or a salt, solvate, or physiologically functional derivative thereof:

wherein:

- 15

D is



- 20 X_1 is hydrogen, C_{1-4} alkyl, C_{1-4} haloalkyl, or C_{1-4} hydroxyalkyl;
 X_2 is hydrogen, C_{1-4} alkyl, C_{1-4} haloalkyl, $C(O)R^1$, or aralkyl;

X_3 is hydrogen or halogen;

X_4 is hydrogen, C_{1-4} alkyl, C_{1-4} haloalkyl, heteroaralkyl, cyanoalkyl, $-(CH_2)_pC=CH(CH_2)_lH$, $-(CH_2)_pC\equiv C(CH_2)_lH$, or C_{3-7} cycloalkyl;

p is 1, 2, or 3;

5 t is 0 or 1;

W is N or C-R, wherein R is hydrogen, halogen, or cyano;

Q_1 is hydrogen, halogen, C_{1-2} haloalkyl, C_{1-2} alkyl, C_{1-2} alkoxy, or C_{1-2} haloalkoxy;

Q_2 is A^1 or A^2 ;

Q_3 is A^1 when Q_2 is A^2 and Q_3 is A^2 when Q_2 is A^1 ;

10 wherein

A^1 is hydrogen, halogen, C_{1-3} alkyl, C_{1-3} haloalkyl, $-OR^1$, and

A^2 is the group defined by $-(Z)_m-(Z^1)-(Z^2)$, wherein

Z is CH_2 and m is 0, 1, 2, or 3, or

Z is NR^2 and m is 0 or 1, or

15 Z is oxygen and m is 0 or 1, or

Z is CH_2NR^2 and m is 0 or 1;

Z^1 is $S(O)_2$, $S(O)$, or $C(O)$; and

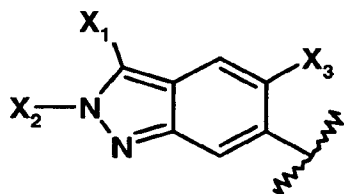
Z^2 is C_{1-4} alkyl, NR^3R^4 , aryl, arylamino, aralkyl, aralkoxy, or heteroaryl,

R^1 is C_{1-4} alkyl;

20 R^2 , R^3 , and R^4 are each independently selected from hydrogen, C_{1-4} alkyl, C_{3-7} cycloalkyl, $-S(O)_2R^5$, and $-C(O)R^5$;

R^5 is C_{1-4} alkyl, or C_{3-7} cycloalkyl; and

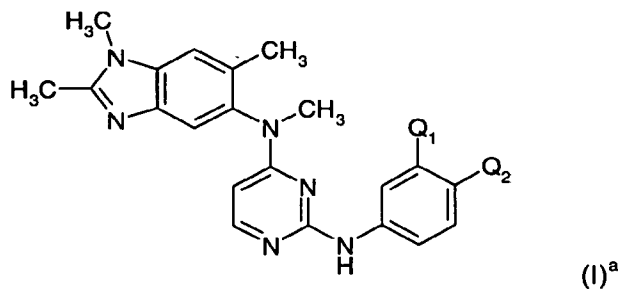
when Z is oxygen then Z^1 is $S(O)_2$ and when D is



25 then X_2 is C_{1-4} alkyl, C_{1-4} haloalkyl, $C(O)R^1$, or aralkyl.

In another embodiment, the compound of formula I is a compound of formula

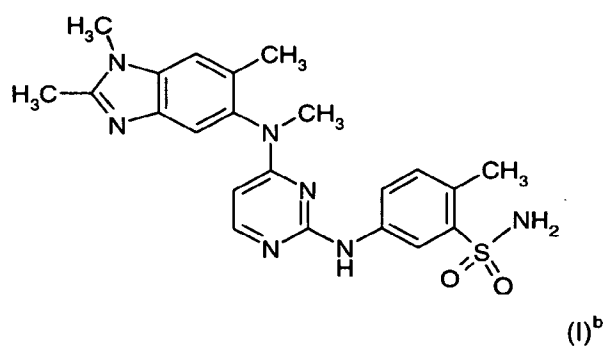
I^a:



or a salt, solvate, or physiologically functional derivative thereof.

In another embodiment, the compound of formula I is a compound of formula

5 I^b:

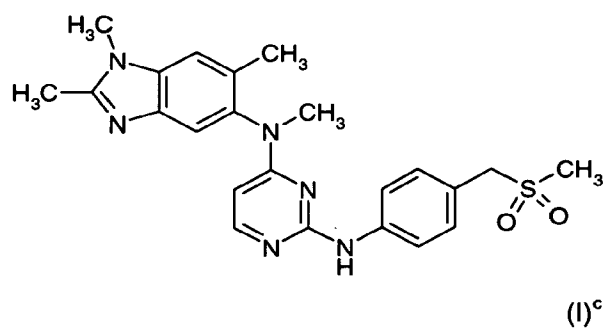


or a salt, solvate or physiological functional derivative thereof.

10

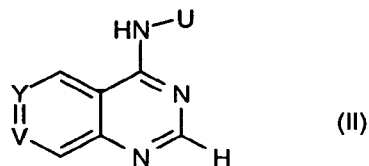
In another embodiment, the compound of formula I is a compound of formula

I^c:



15 or salt, solvate or physiologically functional derivative thereof.

In another embodiment of the present invention is a compound of the formula II:



or a salt, solvate, or physiologically functional derivative thereof;

wherein

- 5 Y is CR¹ and V is N;
or Y is CR¹ and V is CR²;

- R¹ represents a group CH₃SO₂CH₂CH₂NHCH₂-Ar-, wherein Ar is selected from phenyl, furan, thiophene, pyrrole and thiazole, each of which may optionally be substituted by one or two halo, C₁₋₄ alkyl or C₁₋₄ alkoxy groups;
- 10

R² is selected from the group comprising hydrogen, halo, hydroxy, C₁₋₄ alkyl, C₁₋₄ alkoxy, C₁₋₄ alkylamino and di[C₁₋₄ alkyl]amino;

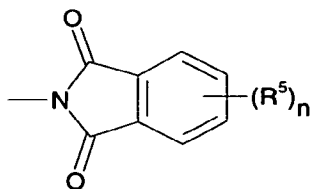
- 15 U represents a phenyl, pyridyl, 3H-imidazolyl, indolyl, isoindolyl, indolinyl, isoindolinyl, 1H-indazolyl, 2,3-dihydro-1H-indazolyl, 1H-benzimidazolyl, 2,3-dihydro-1H-benzimidazolyl or 1H-benzotriazolyl group, substituted by an R³ group and optionally substituted by at least one independently selected R⁴ group;

- 20 R³ is selected from a group comprising benzyl, halo-, dihalo- and trihalobenzyl, benzoyl, pyridylmethyl, pyridylmethoxy, phenoxy, benzyloxy, halo-, dihalo- and trihalobenzyloxy and benzenesulphonyl;

or R³ represents trihalomethylbenzyl or trihalomethylbenzyloxy;

25

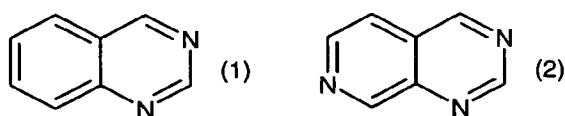
or R³ represents a group of formula



wherein each R⁵ is independently selected from halogen, C₁₋₄ alkyl and C₁₋₄ alkoxy; and n is 0 to 3; and

each R⁴ is independently hydroxy, halogen, C₁₋₄ alkyl, C₂₋₄ alkenyl, C₂₋₄ alkynyl, C₁₋₄ alkoxy, amino, C₁₋₄ alkylamino, di[C₁₋₄ alkyl]amino, C₁₋₄ alkylthio, C₁₋₄ alkylsulphinyl, C₁₋₄ alkylsulphonyl, C₁₋₄ alkylcarbonyl, carboxy, carbamoyl, C₁₋₄ alkoxycarbonyl, C₁₋₄ alkanoylamino, N-(C₁₋₄ alkyl)carbamoyl, N,N-di(C₁₋₄ alkyl)carbamoyl, cyano, nitro and trifluoromethyl.

The definitions for Y and V thus give rise to two possible basic ring systems for the compounds of formula (II). In particular the compounds may contain the following basic ring systems: quinazolines (1) and pyrido-pyrimidines (2):



In a preferred embodiment, the ring system is ring (1).

15

Suitable values for the various groups listed above within the definitions for R¹, R², R⁴ and R⁵ are as follows:

halo is, for example, fluoro, chloro, bromo or iodo; preferably it is fluoro, chloro or bromo, more preferably fluoro or chloro;

20 C₁₋₄ alkyl is, for example, methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl or tert-butyl; preferably it is methyl, ethyl, propyl, isopropyl or butyl, more preferably methyl;

C₂₋₄ alkenyl is, for example, ethenyl, prop-1-enyl or prop-2-enyl; preferably ethenyl;

C₂₋₄ alkynyl is, for example, ethynyl, prop-1-ynyl or prop-2-ynyl; preferably ethynyl;

25 C₁₋₄ alkoxy is, for example, methoxy, ethoxy, n-propoxy, isopropoxy, n-butoxy, isobutoxy, sec-butoxy or tert-butoxy; preferably methoxy, ethoxy, propoxy, isopropoxy or butoxy; more preferably methoxy;

C₁₋₄ alkylamino is, for example, methylamino, ethylamino or propylamino; preferably methylamino;

30 di[C₁₋₄ alkyl]amino is, for example, dimethylamino, diethylamino, N-methyl-N-ethylamino or dipropylamino; preferably dimethylamino;

C₁₋₄ alkylthio is, for example, methylthio, ethylthio, propylthio or isopropylthio, preferably methylthio;

C₁₋₄ alkylsulphinyl is, for example, methylsulphinyl, ethylsulphinyl, propylsulphinyl or isopropylsulphinyl, preferably methylsulphinyl;

C₁₋₄ alkylsulphonyl is, for example, methanesulphonyl, ethylsulphonyl, propylsulphonyl or isopropylsulphonyl, preferably methanesulphonyl;

5 C₁₋₄ alkylcarbonyl is, for example methylcarbonyl, ethylcarbonyl or propylcarbonyl;

C₁₋₄ alkoxy carbonyl is, for example, methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, butoxycarbonyl or tert-butoxycarbonyl;

C₁₋₄ alkanoylamino (where the number of carbon atoms includes the CO functionality) is, for example, formamido, acetamido, propionamido or butyramido;

10 N-(C₁₋₄ alkyl)carbamoyl is, for example, N-methylcarbamoyl or N-ethylcarbamoyl; and N,N-di(C₁₋₄ alkyl)carbamoyl is, for example, N,N-dimethylcarbamoyl, N-methyl-N-ethylcarbamoyl or N,N-diethylcarbamoyl.

15 In a preferred embodiment, Y is CR¹ and V is CR² (ring system (1) above).

In another embodiment, Y is CR¹ and V is N (ring system (2) above).

In one embodiment, R² represents hydrogen or C₁₋₄ alkoxy.

20 In a preferred embodiment, R² represents hydrogen or methoxy.

In another preferred embodiment, R² represents halo; more preferred R² is fluoro.

25 In a preferred embodiment, the group Ar is substituted by one halo, C₁₋₄ alkyl or C₁₋₄ alkoxy group.

In a more preferred embodiment, the group Ar is substituted by a C₁₋₄ alkyl group.

30 In another preferred embodiment, the group Ar does not carry any optional substituents.

35 In a further more preferred embodiment, Ar represents furan, phenyl or thiazole, each of which may optionally be substituted as indicated above.

In a further more preferred embodiment, Ar represents furan or thiazole, each of which may optionally be substituted as indicated above.

5 In a most preferred embodiment, Ar represents unsubstituted furan or thiazole.

The side chain $\text{CH}_3\text{SO}_2\text{CH}_2\text{CH}_2\text{NHCH}_2$ may be linked to any suitable position of the group Ar. Similarly, the group R^1 may be linked to the carbon atom carrying it from any suitable position of the group Ar.

10

In a preferred embodiment, when Ar represents furan the side chain $\text{CH}_3\text{SO}_2\text{CH}_2\text{CH}_2\text{NHCH}_2$ is in the 4-position of the furan ring and the link to the carbon atom carrying the group R^1 is from the 2-position of the furan ring.

15 In another preferred embodiment, when Ar represents furan the side chain $\text{CH}_3\text{SO}_2\text{CH}_2\text{CH}_2\text{NHCH}_2$ is in the 3-position of the furan ring and the link to the carbon atom carrying the group R^1 is from the 2-position of the furan ring.

20 In a most preferred embodiment, when Ar represents furan the side chain $\text{CH}_3\text{SO}_2\text{CH}_2\text{CH}_2\text{NHCH}_2$ is in the 5-position of the furan ring and the link to the carbon atom carrying the group R^1 is from the 2-position of the furan ring.

25 In a further most preferred embodiment, when Ar represents thiazole the side chain $\text{CH}_3\text{SO}_2\text{CH}_2\text{CH}_2\text{NHCH}_2$ is in the 2-position of the thiazole ring and the link to the carbon atom carrying the group R^1 is from the 4-position of the thiazole ring.

30 The R^3 and R^4 groups may be bound to the ring system U by either a carbon atom or a heteroatom of the ring system. The ring system itself may be bound to the bridging NH group by a carbon atom or a heteroatom but is preferably bound by a carbon atom. The R^3 and R^4 groups may be bound to either ring when U represents a bicyclic ring system, but these groups are preferably bound to the ring which is not bound to the bridging NH group in such a case.

In a preferred embodiment U, represents a phenyl, indolyl, or 1H-indazolyl group substituted by an R³ group and optionally substituted by at least one independently selected R⁴ group.

5 In a more preferred embodiment, U represents a phenyl or 1H-indazolyl group substituted by an R³ group and optionally substituted by at least one independently selected R⁴ group.

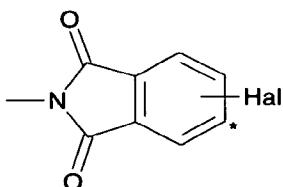
10 In a more preferred embodiment, where U represents a phenyl group the group R³ is in the para- position relative to the bond from U to the linking NH group.

In a further more preferred embodiment, where U represents a 1H-indazolyl group the group R³ is in the 1-position of the indazolyl group.

15 In a preferred embodiment, R³ represents benzyl, pyridylmethyl, phenoxy, benzyloxy, halo-, dihalo- and trihalobenzyloxy and benzenesulphonyl.

In a further preferred embodiment, R³ represents trihalomethylbenzyloxy.

20 In a further preferred embodiment, R³ represents a group of formula



, wherein Hal is Br or Cl, particularly Cl, more especially wherein the Hal substituent is in the position marked with a star in the ring as shown.

25 In a more preferred embodiment, R³ represents benzyloxy, fluorobenzyloxy (especially 3-fluorobenzyloxy), benzyl, phenoxy and benzenesulphonyl.

30 In a further more preferred, embodiment R³ represents bromobenzyloxy (especially 3-bromobenzyloxy) and trifluoromethylbenzyloxy.

In a further preferred embodiment, the ring U is not substituted by an R⁴ group; in an especially preferred embodiment U is phenyl or indazolyl unsubstituted by an R⁴ group.

- 5 In a further preferred embodiment, the ring U is substituted by an R⁴ group selected from halo or C₁₋₄ alkoxy; especially chloro, fluoro or methoxy.

In a more preferred embodiment, the ring U is substituted by an R⁴ group wherein R⁴ represents halo, especially 3-fluoro.

10

In another preferred embodiment, U together with R⁴ represents methoxyphenyl, fluorophenyl, trifluoromethylphenyl or chlorophenyl.

- 15 In a further preferred embodiment, U together with R⁴ represents methoxyphenyl or fluorophenyl.

In another preferred embodiment, the group U together with the substituent(s) R³ and R⁴ represents benzyloxyphenyl, (fluorobenzyloxy)phenyl, (benzenesulphonyl)phenyl, benzylindazolyl or phenoxyphenyl.

20

In still another preferred embodiment, the group U together with the substituent(s) R³ and R⁴ represents benzyloxyphenyl, (3-fluorobenzyloxy)phenyl, (benzenesulphonyl)phenyl or benzylindazolyl.

- 25 In another preferred embodiment, the group U together with the substituent(s) R³ and R⁴ represents (3-bromobenzyloxy)phenyl, (3-trifluoromethylbenzyloxy)phenyl, or (3-fluorobenzyloxy)-3-methoxyphenyl.

- 30 In a more preferred embodiment, the group U together with the substituent(s) R³ and R⁴ represents 3-fluorobenzyloxy-3-chlorophenyl, benzyloxy-3-chlorophenyl, benzyloxy-3-trifluoromethylphenyl, (benzyloxy)-3-fluorophenyl, (3-fluorobenzyloxy)-3-fluorophenyl or (3-fluorobenzyl)indazolyl.

- 35 In another preferred embodiment the group U together with the substituent(s) R³ and R⁴ represents benzyloxyphenyl or (3-fluorobenzyloxy)phenyl.

In a preferred embodiment, there is provided a compound of formula II or a salt, solvate, or physiologically functional derivative thereof wherein V is CR², wherein R² is hydrogen, halo (especially fluoro) or C₁₋₄ alkoxy (especially methoxy); Y is CR¹ wherein R¹ is as defined above in which Ar is unsubstituted phenyl, furan or thiazole; U is phenyl or indazole; R³ is benzyl, fluorobenzyl, benzyloxy, fluorobenzyloxy, bromobenzyloxy, trifluoromethylbenzyloxy, phenoxy or benzenesulphonyl; and R⁴ is not present or is halo (especially chloro or fluoro), or methoxy.

10

In another preferred embodiment, there is provided a compound of formula II or a salt, solvate, or physiologically functional derivative thereof wherein V is CR², wherein R² is hydrogen, halo (especially fluoro) or C₁₋₄ alkoxy (especially methoxy); Y is CR¹ wherein R¹ is as defined above in which Ar is unsubstituted furan or thiazole; U is phenyl; R³ is benzyloxy, fluorobenzyloxy or benzenesulphonyl; and R⁴ is not present or is halo (especially chloro or fluoro), or methoxy.

15

In a preferred embodiment, there is provided a compound of formula II or a salt, solvate, or physiologically functional derivative thereof wherein V is CR², wherein R² is hydrogen, halo (especially fluoro) or C₁₋₄ alkoxy (especially methoxy); Y is CR¹ wherein R¹ is as defined above in which Ar is unsubstituted furan or thiazole; U is indazole; R³ is benzyl or fluorobenzyl; and R⁴ is not present.

20

In a further preferred embodiment, there is provided a compound of formula II or a salt, solvate, or physiologically functional derivative thereof wherein Y is CR², wherein R² is hydrogen, halo (especially fluoro) or C₁₋₄ alkoxy (especially methoxy); V is CR¹ wherein R¹ is as defined above in which Ar is unsubstituted phenyl, furan or thiazole; U is phenyl or indazole; R³ is benzyl, fluorobenzyl, benzyloxy, fluorobenzyloxy, bromobenzyloxy, trifluoromethylbenzyloxy, phenoxy or benzenesulphonyl; and R⁴ is not present or is halo (especially chloro or fluoro), or methoxy.

25

30

In another preferred embodiment, there is provided a compound of formula II or a salt, solvate, or physiologically functional derivative thereof wherein Y is CR², wherein R² is hydrogen, halo (especially fluoro) or C₁₋₄ alkoxy (especially methoxy); V is CR¹ wherein R¹ is as defined above in which Ar is unsubstituted furan or thiazole;

35

U is phenyl; R³ is benzyloxy, fluorobenzyloxy or benzenesulphonyl; and R⁴ is not present or is halo (especially chloro or fluoro), or methoxy.

5 In another preferred embodiment, there is provided a compound of formula II or a salt, solvate, or physiologically functional derivative thereof wherein Y is CR², wherein R² is hydrogen, halo (especially fluoro) or C₁₋₄ alkoxy (especially methoxy); V is CR¹ wherein R¹ is as defined above in which Ar is unsubstituted furan or thiazole; U is indazole; R³ is benzyl or fluorobenzyl; and R⁴ is not present.

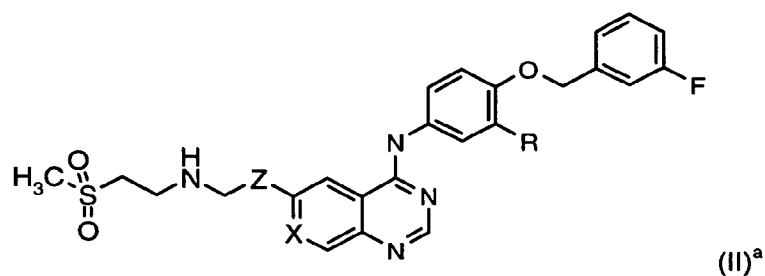
10 In another preferred embodiment, there is provided a compound of formula III or a salt, solvate, or physiologically functional derivative thereof wherein Y is CR², wherein R² is hydrogen, halo (especially fluoro) or C₁₋₄ alkoxy (especially methoxy); V is CR¹ wherein R¹ is as defined above in which Ar is unsubstituted furan or thiazole; U is phenyl; R³ is phenoxy; and R⁴ is not present.

15 In another more preferred embodiment, there is provided a compound of formula II or a salt, solvate, or physiologically functional derivative thereof wherein V is N; Y is CR¹ wherein R¹ is as defined above in which Ar is unsubstituted phenyl, furan or thiazole; U is phenyl or indazole; R³ is benzyl, fluorobenzyl, benzyloxy, fluorobenzyloxy, bromobenzyloxy, trifluoromethylbenzyloxy, phenoxy or
20 benzenesulphonyl; and R⁴ is not present or is halo (especially chloro or fluoro), or methoxy.

In another most preferred embodiment, there is provided a compound of
25 formula II or a salt, solvate, or physiologically functional derivative thereof wherein V is N, Y is CR¹ wherein R¹ is as defined above in which Ar is unsubstituted furan or thiazole; U is phenyl; R³ is benzyloxy, fluorobenzyloxy or benzenesulphonyl; and R⁴ is not present or is halo (especially chloro or fluoro), or methoxy.

30 In another most preferred embodiment, there is provided a compound of formula II or a salt, solvate, or physiologically functional derivative thereof wherein V is N, Y is CR¹ wherein R¹ is as defined above in which Ar is unsubstituted furan or thiazole; U is indazole; R³ is benzyl or fluorobenzyl; and R⁴ is not present.

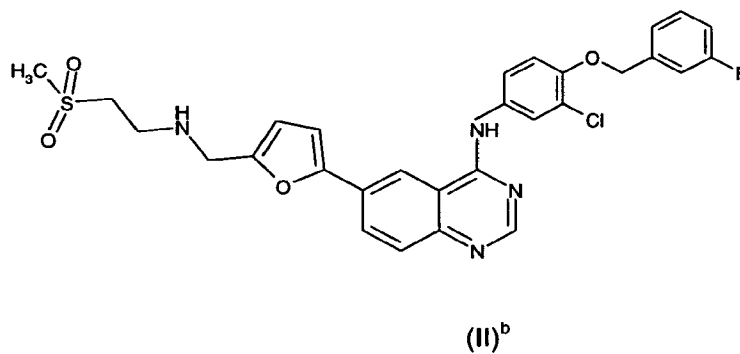
In another most preferred embodiment, the compound of formula II is a compound of formula II^a:



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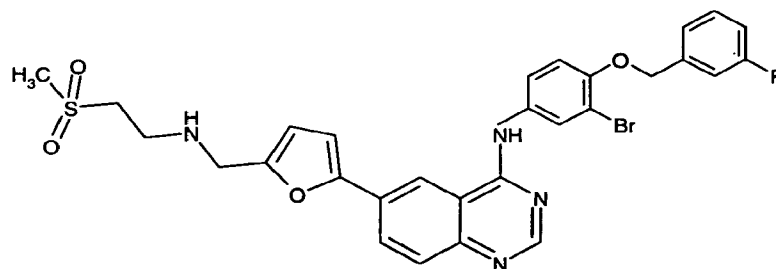
or a salt, solvate or physiologically functional derivative thereof, wherein R is -Cl or -Br, X is CH, N, or CF, and Z is thiazole or furan.

In another most preferred embodiment, the compound of formula II is a compound of formula II^b:



15

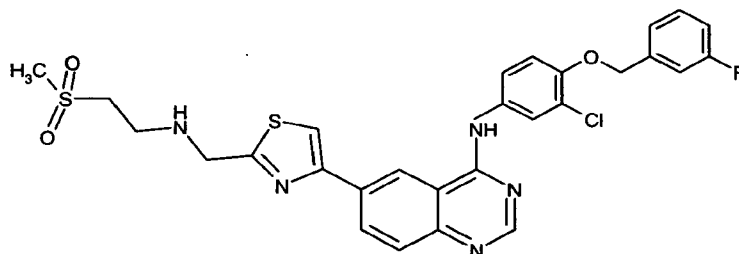
In another most preferred embodiment, the compound of formula II is a compound of formula II^c:



(II)^c

or a salt, solvate or physiologically functional derivative thereof.

In another most preferred embodiment, the compound of formula II is a
5 compound of formula II^d:

(II)^d

or a salt, solvate or physiologically functional derivative thereof.

10 In a most preferred embodiment the compound of formula I is a compound of formula I^b or a salt, solvate or physiologically functional derivative thereof, and the compound of formula II is a compound of formula II^b or a salt, solvate or physiologically functional derivative thereof.

15 In a most preferred embodiment the compound of formula I is a monohydrochloride salt of a compound of formula I^b, and the compound of formula II is a monohydrate ditosylate salt of a compound of the formula II^b.

20 In another most preferred embodiment the compound of formula I is a monohydrochloride salt of a compound of formula I^b, and the compound of formula II is an anhydrous ditosylate salt of a compound of the formula II^b.

25 In a preferred embodiment, the cancer treatment method of the claimed invention includes a compound of formula I or a salt, solvate or physiologically functional derivative thereof and a compound of formula II or a salt, solvate or physiologically functional derivative thereof.

30 In the method of this invention, the compound of formula I or salt, solvate, or physiologically functional derivative thereof and the compound of formula II or salt, solvate, or physiologically functional derivative thereof may be employed in combination concomitantly or sequentially in any therapeutically appropriate

combination. The compounds may be employed in combination in accordance with the invention by administration concomitantly in (1) a unitary pharmaceutical composition including both compounds or (2) separate pharmaceutical compositions each including one of the compounds. Alternatively, the compounds may be
5 administered separately in a sequential manner wherein one is administered first and the other second or vice versa. Such sequential administration may be close in time or remote in time.

The cancer treatment method of the present invention may also include
10 administration of at least one additional cancer treatment therapy in combination concomitantly or sequentially in any therapeutically appropriate combination with the combinations of the present invention. The additional cancer treatment therapy may include radiation therapy, surgical therapy and/or at least one additional chemotherapeutic therapy including administration of at least one additional anti-
15 neoplastic agent.

In a preferred embodiment, the cancer treated by the method of the invention is breast, non-small cell lung, prostate, colorectal, renal, or bladder cancer. In another preferred embodiment, the cancer treated by the method of the invention is
20 mesothelioma, hepatobiliary cancer, multiple myeloma, sarcoma, or leukemia.

While it is possible that, for use in therapy, the compound of formula I or a salt, solvate or physiologically function derivative thereof, or the compound of formula II or a salt, solvate or physiologically function derivative thereof, may be
25 administered as the raw chemical, it is possible to present the active ingredient as a pharmaceutical composition. As indicated above, such elements of the pharmaceutical combination utilized may be presented in separate pharmaceutical compositions or formulated together in one pharmaceutical formulation. Accordingly, the invention further provides a combination of pharmaceutical compositions one of
30 which includes a compound of the formula I or a salt, solvate, or physiologically functional derivative thereof, and one or more pharmaceutically acceptable carriers, diluents, or excipients and a pharmaceutical composition containing a compound of the formula II or a salt, solvate, or physiologically functional derivative thereof, and one or more pharmaceutically acceptable carriers, diluents, or excipients.

35

Alternatively, a pharmaceutical composition is provided which includes a compound of the formula I or a salt, solvate, or physiologically functional derivative thereof, a compound of the formula II or a salt, solvate, or physiologically functional derivative thereof and one or more pharmaceutically acceptable carriers, diluents, or excipients . The compound of formula I or a salt, solvate, or physiologically functional derivative thereof, and the compound of formula II or a salt, solvate, or physiologically functional derivative thereof are as described above and may be utilized in any of the combinations described above in the method of treating cancer of the present invention. A preferred composition may further comprise the preferred compounds, as described above, and one or more pharmaceutically acceptable carriers, diluents, or excipients.

The carrier(s), diluent(s) or excipient(s) must be acceptable in the sense of being compatible with the other ingredients of the formulation and not deleterious to the recipient thereof. According to another aspect of the invention there is also provided a process for the preparation of a pharmaceutical formulation including admixing a compound of the formula I or a salt, solvate, or physiologically functional derivative thereof, and/or a compound of the formula II or a salt, solvate, or physiologically functional derivative thereof, with one or more pharmaceutically acceptable carriers, diluents or excipients.

The components of the pharmaceutical compositions, of the present invention, may be formulated for administration by any route, and the appropriate route will depend on the specific cancer being treated as well as the subjects to be treated. Suitable pharmaceutical formulations include those for oral, rectal, nasal, topical (including buccal, sub-lingual, and transdermal), vaginal or parenteral (including intramuscular, sub-cutaneous, intravenous, and directly into the affected tissue) administration or in a form suitable for administration by inhalation or insufflation. The formulations may, where appropriate, be conveniently presented in discrete dosage units and may be prepared by any of the methods well known in the pharmacy art.

Pharmaceutical formulations adapted for oral administration may be presented as discrete units such as capsules or tablets; powders or granules; solutions or suspensions in aqueous or non-aqueous liquids; edible foams or whips; or oil-in-water liquid emulsions or water-in-oil liquid emulsions.

For instance, for oral administration in the form of a tablet or capsule, the active drug component can be combined with an oral, non-toxic pharmaceutically acceptable inert carrier such as ethanol, glycerol, water and the like. Powders are prepared by comminuting the compound to a suitable fine size and mixing with a similarly comminuted pharmaceutical carrier such as an edible carbohydrate, as, for example, starch or mannitol. Flavoring, preservative, dispersing and coloring agents can also be present.

Capsules are made by preparing a powder mixture as described above, and filling formed gelatin sheaths. Glidants and lubricants such as colloidal silica, talc, magnesium stearate, calcium stearate or solid polyethylene glycol can be added to the powder mixture before the filling operation. A disintegrating or solubilizing agent such as agar-agar, calcium carbonate or sodium carbonate can also be added to improve the availability of the medicament when the capsule is ingested.

Moreover, when desired or necessary, suitable binders, lubricants, disintegrating agents and coloring agents can also be incorporated into the mixture. Suitable binders include starch, gelatin, natural sugars such as glucose or beta-lactose, corn sweeteners, natural and synthetic gums such as acacia, tragacanth or sodium alginate, carboxymethylcellulose, polyethylene glycol, waxes and the like. Lubricants used in these dosage forms include sodium oleate, sodium stearate, magnesium stearate, sodium benzoate, sodium acetate, sodium chloride and the like. Disintegrators include, without limitation, starch, methylcellulose, agar, bentonite, xanthan gum and the like. Tablets are formulated, for example, by preparing a powder mixture, granulating or slugging, adding a lubricant and disintegrant and pressing into tablets. A powder mixture is prepared by mixing the compound, suitably comminuted, with a diluent or base as described above, and optionally, with a binder such as carboxymethylcellulose, an aliginat, gelatin, or polyvinyl pyrrolidone, a solution retardant such as paraffin, a resorption accelerator such as a quaternary salt and/or an absorption agent such as bentonite, kaolin or dicalcium phosphate. The powder mixture can be granulated by wetting with a binder such as syrup, starch paste, acadia mucilage or solutions of cellulosic or polymeric materials and forcing through a screen. As an alternative to granulating, the powder mixture can be run through the tablet machine and the result is imperfectly formed slugs broken into granules. The granules can be lubricated to prevent sticking to the

tablet forming dies by means of the addition of stearic acid, a stearate salt, talc or mineral oil. The lubricated mixture is then compressed into tablets. The compounds of the present invention can also be combined with a free flowing inert carrier and compressed into tablets directly without going through the granulating or slugging steps. A clear or opaque protective coating consisting of a sealing coat of shellac, a coating of sugar or polymeric material and a polish coating of wax can be provided. Dyestuffs can be added to these coatings to distinguish different unit dosages.

Oral fluids such as solution, syrups and elixirs can be prepared in dosage unit form so that a given quantity contains a predetermined amount of the compound. Syrups can be prepared by dissolving the compound in a suitably flavored aqueous solution, while elixirs are prepared through the use of a non-toxic alcoholic vehicle. Suspensions can be formulated by dispersing the compound in a non-toxic vehicle. Solubilizers and emulsifiers such as ethoxylated isostearyl alcohols and polyoxy ethylene sorbitol ethers, preservatives, flavor additive such as peppermint oil or natural sweeteners or saccharin or other artificial sweeteners, and the like can also be added.

Where appropriate, dosage unit formulations for oral administration can be microencapsulated. The formulation can also be prepared to prolong or sustain the release as for example by coating or embedding particulate material in polymers, wax or the like.

The components of the pharmaceutical compositions of the present invention can also be administered in the form of liposome delivery systems, such as small unilamellar vesicles, large unilamellar vesicles and multilamellar vesicles. Liposomes can be formed from a variety of phospholipids, such as cholesterol, stearylamine or phosphatidylcholines.

The components of the pharmaceutical compositions of the present invention may also be delivered by the use of monoclonal antibodies as individual carriers to which the compound molecules are coupled. The compounds may also be coupled with soluble polymers as targetable drug carriers. Such polymers can include polyvinylpyrrolidone, pyran copolymer, polyhydroxypropylmethacrylamide-phenol, polyhydroxyethylaspartamidophenol, or polyethyleneoxidepolylysine substituted with palmitoyl residues. Furthermore, the compounds may be coupled to a class of

biodegradable polymers useful in achieving controlled release of a drug, for example, polylactic acid, polyepsilon caprolactone, polyhydroxy butyric acid, polyorthoesters, polyacetals, polydihydropyrans, polycyanoacrylates and cross-linked or amphipathic block copolymers of hydrogels.

5

Pharmaceutical formulations adapted for transdermal administration may be presented as discrete patches intended to remain in intimate contact with the epidermis of the recipient for a prolonged period of time. For example, the active ingredient may be delivered from the patch by iontophoresis as generally described in Pharmaceutical Research, 3(6), 318 (1986).

10

Pharmaceutical formulations adapted for topical administration may be formulated as ointments, creams, suspensions, lotions, powders, solutions, pastes, gels, sprays, aerosols or oils.

15

For treatments of the eye or other external tissues, for example mouth and skin, the formulations are preferably applied as a topical ointment or cream. When formulated in an ointment, the active ingredient may be employed with either a paraffinic or a water-miscible ointment base. Alternatively, the active ingredient may be formulated in a cream with an oil-in-water cream base or a water-in-oil base.

20

Pharmaceutical formulations adapted for topical administrations to the eye include eye drops wherein the active ingredient is dissolved or suspended in a suitable carrier, especially an aqueous solvent.

25

Pharmaceutical formulations adapted for topical administration in the mouth include lozenges, pastilles and mouthwashes.

Pharmaceutical formulations adapted for rectal administration may be presented as suppositories or as enemas.

30

Pharmaceutical formulations adapted for nasal administration wherein the carrier is a solid include a coarse powder having a particle size for example in the range 20 to 500 microns which is administered in the manner in which snuff is taken, i.e. by rapid inhalation through the nasal passage from a container of the powder held close up to the nose. Suitable formulations wherein the carrier is a liquid, for

35

administration as a nasal spray or as nasal drops, include aqueous or oil solutions of the active ingredient.

Pharmaceutical formulations adapted for administration by inhalation include
5 fine particle dusts or mists, which may be generated by means of various types of metered, dose pressurised aerosols, nebulizers or insufflators.

Pharmaceutical formulations adapted for vaginal administration may be presented as pessaries, tampons, creams, gels, pastes, foams or spray
10 formulations.

Pharmaceutical formulations adapted for parenteral administration include aqueous and non-aqueous sterile injection solutions which may contain anti-oxidants, buffers, bacteriostats and solutes which render the formulation isotonic with
15 the blood of the intended recipient; and aqueous and non-aqueous sterile suspensions which may include suspending agents and thickening agents. The formulations may be presented in unit-dose or multi-dose containers, for example sealed ampoules and vials, and may be stored in a freeze-dried (lyophilized) condition requiring only the addition of the sterile liquid carrier, for example water for
20 injections, immediately prior to use. Extemporaneous injection solutions and suspensions may be prepared from sterile powders, granules and tablets.

It should be understood that in addition to the ingredients particularly mentioned above, the formulations may include other agents conventional in the art
25 having regard to the type of formulation in question, for example those suitable for oral administration may include flavouring agents.

A therapeutically effective amount of the components of the pharmaceutical compositions of the present invention will depend on a number of factors including,
30 but not limited to, the age and weight of the mammal, the precise disorder requiring treatment and its severity, the nature of the formulation, and the route of administration, and will ultimately be at the discretion of the attendant physician or veterinarian. Typically, the components of the pharmaceutical compositions of the present invention will be given for treatment in the range of 0.1 to 100 mg/kg body
35 weight of recipient (mammal) per day and more usually in the range of 1 to 10 mg/kg

body weight per day. Acceptable daily dosages, may be from about 0.1 to about 1000 mg/day, and preferably from about 0.1 to about 100 mg/day.

5 The pharmaceutical compositions, including compounds of formula I or salts, solvates, and physiologically functional derivatives thereof and compounds of formula II or salts, solvates, and physiologically functional derivatives thereof, described above, are useful in therapy and in the preparation of medicaments for treating cancer in a mammal.

10 In one embodiment, the mammal in the methods and uses of the present invention is a human.

The following examples are intended for illustration only and are not intended to limit the scope of the invention in any way. The physical data given for the
15 compounds exemplified is consistent with the assigned structure of those compounds.

EXAMPLES

As used herein the symbols and conventions used in these processes,
20 schemes and examples are consistent with those used in the contemporary scientific literature, for example, the *Journal of the American Chemical Society* or the *Journal of Biological Chemistry*. Standard single-letter or three-letter abbreviations are generally used to designate amino acid residues, which are assumed to be in the L-configuration unless otherwise noted. Unless otherwise noted, all starting materials
25 were obtained from commercial suppliers and used without further purification. Specifically, the following abbreviations may be used in the examples and throughout the specification:

g (grams);	mg (milligrams);
L (liters);	mL (milliliters);
30 μ L (microliters);	psi (pounds per square inch);
M (molar);	mM (millimolar);
i. v. (intravenous);	Hz (Hertz);
MHz (megahertz);	mol (moles);
mmol (millimoles);	RT (room temperature);
35 min (minutes);	h (hours);

	mp (melting point);	TLC (thin layer chromatography);
	T _r (retention time);	RP (reverse phase);
	MeOH (methanol);	<i>i</i> -PrOH (isopropanol);
	TEA (triethylamine);	TFA (trifluoroacetic acid);
5	TFAA (trifluoroacetic anhydride);	THF (tetrahydrofuran);
	DMSO (dimethylsulfoxide);	EtOAc (ethyl acetate);
	DME (1,2-dimethoxyethane);	DCM (dichloromethane);
	DCE (dichloroethane);	DMF (<i>N,N</i> -dimethylformamide);
	DMPU (<i>N,N'</i> -dimethylpropyleneurea);	CDI (1,1-carbonyldiimidazole);
10	IBCF (isobutyl chloroformate);	HOAc (acetic acid);
	HOSu (<i>N</i> -hydroxysuccinimide);	HOBT (1-hydroxybenzotriazole);
	mCPBA (meta-chloroperbenzoic acid);	Et (ethyl);
	EDC (ethylcarbodiimide hydrochloride);	BOC (<i>tert</i> -butoxycarbonyl);
	Fmoc (9-fluorenylmethoxycarbonyl);	DCC (dicyclohexylcarbodiimide);
15	CBZ (benzyloxycarbonyl);	Ac (acetyl);
	atm (atmosphere);	TMSE (2-(trimethylsilyl)ethyl);
	TMS (trimethylsilyl);	TIPS (triisopropylsilyl);
	TBS (<i>t</i> -butyldimethylsilyl);	OMe (methoxy);
	DMAP (4-dimethylaminopyridine);	Me (methyl);
20	HPLC (high pressure liquid chromatography);	
	BOP (bis(2-oxo-3-oxazolidinyl)phosphinic chloride);	
	TBAF (tetra- <i>n</i> -butylammonium fluoride);	tBu (tert-butyl).

25 All references to ether are to diethyl ether; brine refers to a saturated aqueous solution of NaCl. Unless otherwise indicated, all temperatures are expressed in °C (degrees Centigrade). All reactions conducted under an inert atmosphere at room temperature unless otherwise noted.

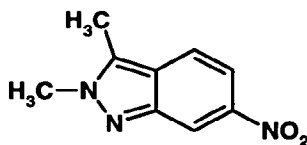
30 For the compounds of the formula I, ¹H NMR spectra were recorded on a Varian VXR-300, a Varian Unity-300, a Varian Unity-400 instrument, or a General Electric QE-300. Chemical shifts are expressed in parts per million (ppm, δ units). Coupling constants are in units of hertz (Hz). Splitting patterns describe apparent multiplicities and are designated as s (singlet), d (doublet), t (triplet), q (quartet), m (multiplet), br (broad).

Low-resolution mass spectra (MS) were recorded on a JOEL JMS-AX505HA, JOEL SX-102, or a SCIEX-APIiii spectrometer; high resolution MS were obtained using a JOEL SX-102A spectrometer. All mass spectra were taken under electrospray ionization (ESI), chemical ionization (CI), electron impact (EI) or by fast atom bombardment (FAB) methods. Infrared (IR) spectra were obtained on a Nicolet 510 FT-IR spectrometer using a 1-mm NaCl cell. All reactions were monitored by thin-layer chromatography on 0.25 mm E. Merck silica gel plates (60F-254), visualized with UV light, 5% ethanolic phosphomolybdic acid or p-anisaldehyde solution. Flash column chromatography was performed on silica gel (230-400 mesh, Merck). Optical rotations were obtained using a Perkin Elmer Model 241 Polarimeter. Melting points were determined using a Mel-Temp II apparatus and are uncorrected.

For the compounds of the formula II ^1H NMR spectra were obtained at 500MHz on a Bruker AMX500 spectrophotometer, on a Bruker spectrophotometer at 300MHz, on a Bruker AC250 or Bruker AM250 spectrophotometer at 250MHz and on a Varian Unity Plus NMR spectrophotometer at 300 or 400 MHz. J values are given in Hz. Mass spectra were obtained on one of the following machines: VG Micromass Platform (electrospray positive or negative), HP5989A Engine (thermospray positive) or Finnigan-MAT LCQ (ion trap) mass spectrometer. Analytical thin layer chromatography (tlc) was used to verify the purity of some intermediates which could not be isolated or which were too unstable for full characterization, and to follow the progress of reactions. Unless otherwise stated, this was done using silica gel (Merck Silica Gel 60 F254).

The free base, HCl salts, and ditosylate salts of the compound of formula (II) may be prepared according to the procedures of the International Patent Application No. PCT/EP99/00048, filed January 8, 1999, and published as WO 99/35146 on July 15. Such application is incorporated herein by reference to the extent it teaches the preparation of the compound of formula (II) and salts thereof. Some of such procedures are recited again herein as well as additional variations and procedures.

The following examples describe the syntheses of intermediates particularly useful in the synthesis of compounds of formula (I):

Intermediate Example 1**Preparation of 2,3-dimethyl-6-nitro-2H-indazole****Procedure 1:**

- 5 To a stirred solution of 18.5 g (0.11 mol) of 3-methyl-6-nitro-1H-indazole in 350 ml acetone, at room temperature, was added 20 g (0.14 mol) of trimethyloxonium tetrafluoroborate. After the solution was allowed to stir under argon for 3 hours, the solvent was removed under reduced pressure. To the resulting solid was added saturated aqueous NaHCO₃ (600 mL) and a 4:1 mixture of
- 10 chloroform-isopropanol (200 ml), the mixture was agitated and the layers were separated. The aqueous phase was washed with additional chloroform: isopropanol (4 x 200 mL) and the combined organic phase was dried (Na₂SO₄). Filtration and removal of solvent gave a tan solid. The solid was washed with ether (200 mL) to afford 2,3-dimethyl-6-nitro-2H-indazole as a yellow solid (15.85 g, 73 %). ¹H NMR
- 15 (300 MHz, DMSO-d₆) δ 8.51 (s, 1H), 7.94 (d, *J* = 9.1 Hz, 1H), 7.73 (d, *J* = 8.9 Hz, 1H), 4.14 (s, 3H), 2.67 (s, 3H). MS (ES+, *m/z*) 192 (M+H).

Procedure 2:

- 20 Trimethyl orthoformate (11 mmol, 1.17 g) was added over a 2 min period to a solution of boron trifluoride etherate (12.5 mmol, 1.77 g in methylene chloride (2.0 mL) which had been cooled to -30 °C. The mixture was warmed to 0 °C for 15 min and was then cooled to -70 °C. The nitro indazole (10 mmol, 1.77 g) was slurried in methylene chloride (30 mL) and was added all at once to the cooled mixture. The mixture was stirred at -70 °C for 15 min and at ambient temperature for 17 h. After
- 25 17 h the mixture was red and heterogeneous. The reaction mixture was quenched with saturated sodium bicarbonate solution (20 mL) and the organic layer separated. The aqueous layer was extracted with methylene chloride (30 mL). The methylene chloride layers were combined and extracted with water (30 mL). The methylene chloride layer was distilled under reduced pressure until ~ 10 mL remained.
- 30 Propanol (10 mL) was added and the remainder of the methylene chloride removed under reduced pressure, resulting in a yellow slurry. The product was isolated by filtration to give 2,3-dimethyl-6-nitro-2H-indazole (65 %, 7mmol, 1.25 g) as a light

yellow powder. ^1H NMR (300 MHz, DMSO-d_6) δ 8.51 (s, 1H), 7.94 (d, J = 9.1 Hz, 1H), 7.73 (d, J = 8.9 Hz, 1H), 4.14 (s, 3H), 2.67 (s, 3H). MS (ES+, m/z) 192 (M+H).

Procedure 3:

5 In a 25 ml round bottom flask 3-methyl-6-nitroindazole (7.27 mmol, 1.28 g) was dissolved with stirring in DMSO (4.0 mL) and was treated with concentrated sulfuric acid (7.27 mmol, 0.73 g) to yield a thick slurry. The slurry was treated with dimethyl sulfate (21.1 mmol, 2.66 g). The mixture was heated under nitrogen at 50 °C for 72 h. After 72 h a thick yellow slurry was obtained. The slurry was cooled and
10 was slowly treated with saturated sodium bicarbonate solution (10 mL). The mixture was extracted with methylene chloride (2 x 20 mL). The methylene chloride layers were combined and back extracted with water (20 mL). The methylene chloride layer was treated with propanol (10 mL) and the methylene chloride was removed by distillation under reduced pressure. The solid was isolated by filtration and the
15 yellow solid washed with heptane (5 mL) and air-dried. The 2,3-dimethyl-6-nitro-2H-indazole product (70%, 0.97 g) was obtained as a light yellow solid. ^1H NMR (300 MHz, DMSO-d_6) δ 8.51 (s, 1H), 7.94 (d, J = 9.1 Hz, 1H), 7.73 (d, J = 8.9 Hz, 1H), 4.14 (s, 3H), 2.67 (s, 3H). MS (ES+, m/z) 192 (M+H).

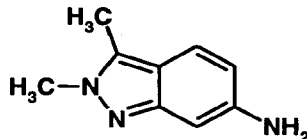
20 **Procedure 4:**

 Into a 250 mL 3-necked round bottom flask was placed 3-methyl-6-nitro-1H-indazole sulfuric acid salt (5.0 g, 18.2 mmol) and methylene chloride (25 mL). The mixture was stirred at 25 °C and was treated with DMSO (5 mL). Dimethyl sulfate
25 (6.7 g, 5.0 mL, 53.0 mmol) was added via syringe and the reaction was heated at reflux in a 70 °C bath. After 7 h HPLC analysis showed 9% starting material. At this point heating was stopped and the workup begun. Saturated sodium bicarbonate solution (35 mL) was added to the reaction mixture at RT. The layers were allowed to separate and the aqueous layer was extracted with methylene chloride (25 mL).
30 The methylene chloride layers were combined and washed with water (2 x 25 mL). The methylene chloride layer was distilled under reduced pressure until half the volume was removed. Propanol (25 mL) was added and distillation under reduced pressure was continued until all the methylene chloride had been removed. This yielded a yellow slurry, which was allowed to stir at 25 °C for 1 h. The product was
35 isolated via filtration and the resulting yellow solid was washed with heptane (10 mL). This yielded 2,3-dimethyl-6-nitro-2H-indazole (70%, 2.43 g) as a yellow solid. ^1H

NMR (300 MHz, DMSO- d_6) δ 8.51 (s, 1H), 7.94 (d, J = 9.1 Hz, 1H), 7.73 (d, J = 8.9 Hz, 1H), 4.14 (s, 3H), 2.67 (s, 3H). MS (ES+, m/z) 192 (M+H).

5 Intermediate Example 2

Preparation of 2,3-dimethyl-6-amino-2H-indazole



Procedure 1:

10 To a stirred solution of 2,3-dimethyl-6-nitro-2H-indazole (1.13 g) in 2-methoxyethyl ether (12 ml), at 0 °C, was added a solution of 4.48 g of tin(II) chloride in 8.9 ml of concentrated HCl dropwise over 5 min. After the addition was complete, the ice bath was removed and the solution was allowed to stir for an additional 30 min. Approximately 40 ml of diethyl ether was added to reaction, resulting in
15 precipitate formation. The resulting precipitate was isolated by filtration and washed with diethyl ether, and afforded a yellow solid (1.1 g, 95 %), the HCl salt 2,3-dimethyl-2H-indazol-6-amine. ^1H NMR (300 MHz, DMSO- d_6) δ 7.77 (d, J = 8.9 Hz, 1H), 7.18 (s, 1H), 7.88 (m, 1H), 4.04 (s, 3H), 2.61 (s, 3H). MS (ES+, m/z) 162 (M+H).

20

Procedure 2:

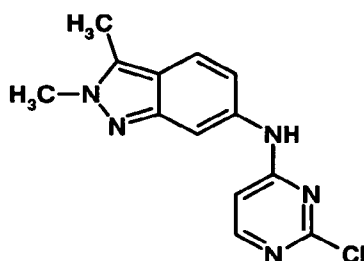
A 2-L 3-necked round bottom flask was fitted with nitrogen inlet and outlet and with mechanical stirring. A moderate nitrogen flow was initiated and the reactor was charged with 10 % Pd/C (50% water wet, 6.0 g). Stirring was initiated and the
25 reactor was charged with methanol (750 mL) and the product of Intermediate Example 1 (50 g). Ammonium formate (82.54 g) was dissolved in water (120 mL). The water solution of ammonium formate was added to the reaction solution at an addition rate, which kept the reaction temperature at or between 25 and 30 °C. The reaction was allowed to proceed at 25 °C. After 6 h the reaction was judged to be
30 finished based on HPLC analysis. The mixture was filtered and the catalyst washed with methanol (50 mL). The methanol layers were combined and the solvent removed under reduced pressure. The residue was dissolved in water (200 mL) and

was extracted with methylene chloride (3 x 250 mL). The methylene chloride layers were combined and solvent removed under vacuum to remove approximately half the solvent. Heptane (400 mL) was added and the vacuum distillation continued until approximately 300 mL reaction product slurry remained. The product was isolated by filtration and dried under vacuum at 50 °C for 4 h. to yield 2,3-dimethyl-6-amino-2H-indazole as the free base. (40.76 g, 96.7 %). ¹H NMR (300 MHz, DMSO-d₆) δ 7.31 (d, *J* = 8.9 Hz, 1H), 6.45 (d, *J* = 8.9 Hz, 1H), 6.38 (s, 1H), 4.95 (s, br, 2H), 3.85 (s, 3H), 2.44 (s, 3H) MS (ES+, *m/z*) 162 (M+H).

10

Intermediate Example 3

Preparation of N-(2-chloropyrimidin-4-yl)-2,3-dimethyl-2H-indazol-6-amine



Procedure 1

To a stirred solution of the product of Intermediate Example 2 (2.97 g, .015 mol) and NaHCO₃ (5.05 g, .06 mol) in THF (15 mL) and ethanol (60 mL) was added 2,4-dichloropyrimidine (6.70 g, .045 mol) at rt. After the reaction was stirred for four hours at 85 °C, the suspension was cooled to rt., filtered and washed thoroughly with ethyl acetate. The filtrate was concentrated under reduced pressure, and the resulting solid was triturated with ethyl acetate to yield *N*-(2-chloropyrimidin-4-yl)-2,3-dimethyl-2H-indazol-6-amine (89 %, 3.84 g). ¹H NMR (400 MHz, DMSO-d₆) δ 7.28 (d, *J* = 9.0 Hz, 1H), 6.42 (d, *J* = 8.8 Hz, 1H), 6.37 (s, 1H), 5.18 (br s, 1H), 3.84 (s, 3H), 2.43 (s, 3H). MS (ES+, *m/z*) 274 (M+H).

25

Procedure 2

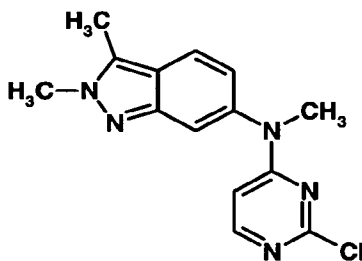
To a 1-L 3-necked flask equipped with air-driven mechanical stirrer, thermometer, and nitrogen inlet/outlet was charged a solution of the product of Intermediate Example 2 (32.89 g, 0.204 mol, 1.0 equiv) in 425 mL (13 volumes) of EtOH/THF (4/1), sodium bicarbonate (51.42 g, 0.612 mol, 3.0 equiv) and then 2,4-

dichloropyrimidine (45.59 g, 0.306 mol, 1.5 equiv). The flask contents were heated to 75 °C and held at 74 – 76 °C for 6 – 7 hrs. The progress of the reaction was checked by HPLC (the product of Intermediate Example 2 < 2%). The reaction contents were cooled to 20 – 25 °C over 30 min, and kept at 20 – 25 °C for 30 min.

- 5 Then the reaction contents were further cooled to 10 - 12 °C over 30 min, and kept at that temperature for an additional 10 min. The contents were filtered and filter cake washed with EtOAc (2 x 100 mL, 3.0 volumes), and deionized water (514 mL, 15.6 volumes). The filter cake was then dried in a vacuum oven at 35 °C overnight to afford the desired product 44.75 g as a white solid (80.1%). ¹H NMR (400 MHz, DMSO-d₆) δ 7.28 (d, *J* = 9.0 Hz, 1H), 6.42 (d, *J* = 8.8 Hz, 1H), 6.37 (s, 1H), 5.18 (br s, 1H), 3.84 (s, 3H), 2.43 (s, 3H). MS (ES⁺, *m/z*) 274 (M+H).
- 10

Intermediate Example 4

- 15 *Preparation of N-(2-chloropyrimidin-4-yl)-N,2,3-trimethyl-2H-indazol-6-amine*



Procedure 1

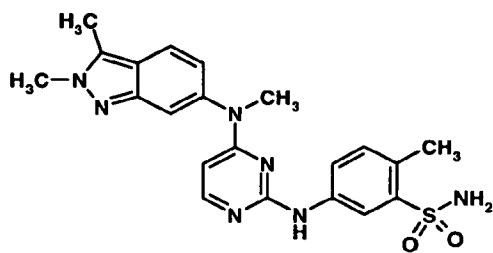
- To a stirred solution of the product of Intermediate Example 3 (7.37 g) in DMF (50 ml) was added Cs₂CO₃ (7.44 g, 2 eqv.) and iodomethane (1.84 ml, 1.1 eqv.) at room temperature. The mixture was stirred at rt overnight. The reaction mixture was then poured into an ice-water bath, and the precipitate was collected via filtration and washed with water. The precipitate was air-dried to afford *N*-(2-chloropyrimidin-4-yl)-*N*,2,3-trimethyl-2*H*-indazol-6-amine as an off-white solid (6.43 g, 83%). ¹H NMR (400 MHz, DMSO-d₆) δ 7.94 (d, *J* = 6.0 Hz, 1H), 7.80 (d, *J* = 7.0 Hz, 1H), 7.50 (d, *J* = 1.0 Hz, 1H), 6.88 (m, 1H), 6.24 (d, *J* = 6.2 Hz, 1H), 4.06 (s, 3H), 3.42 (s, 3H), 2.62 (s, 3H). MS (ES⁺, *m/z*) 288 (M+H).
- 20
- 25

Procedure 2

A 3L 3-necked flask equipped with air-driven mechanical stirrer, thermometer, addition funnel and nitrogen inlet/outlet was charged with DMF (272 mL, 5 volumes) and the product of Intermediate Example 3 (54.4 g, 0.20 mol, 1.0 equiv) with stirring. The reaction mixture was further charged with cesium carbonate (194.5 g, 0.60 mol, 3.0 equiv) while maintaining the reaction temperature between 20 ~ 25 °C. The reaction mixture was stirred at 20 ~ 25 °C for 10 minutes. Iodomethane (45.1 g, 0.32 mol, 1.6 equiv) was charged over ~ 10 minutes while maintaining the temperature 20 ~ 30°C. The reaction mixture was stirred at 20 ~ 30 °C (Typically, the reaction is complete in 1 ~ 2 hours). Deionized H₂O (925 mL, 17 volumes) was added over ~ 30 minutes while maintaining the temperature at 25 ~ 40 °C. The reaction mixture was stirred at 20 ~ 25 °C for 40 minutes. The product was isolated by filtration and then the filter cake washed with H₂O / DMF (6 : 1, 252 mL, 4.6 volumes). The wet cake was dried under vacuum at 40 ~ 45 °C and *N*-(2-chloropyrimidin-4-yl)-*N*,2,3-trimethyl-2*H*-indazol-6-amine (51.7 g, 90.4%) was isolated as a yellow solid. ¹H NMR (400 MHz, DMSO-*d*₆) δ 7.94 (d, *J* = 6.0 Hz, 1H), 7.80 (d, *J* = 7.0 Hz, 1H), 7.50 (d, *J* = 1.0 Hz, 1H), 6.88 (m, 1H), 6.24 (d, *J* = 6.2 Hz, 1H), 4.06 (s, 3H), 3.42 (s, 3H), 2.62 (s, 3H). MS (ES+, *m/z*) 288 (M+H).

Example 1

5-({4-[(2,3-dimethyl-2*H*-indazol-6-yl)(methyl)amino]pyrimidin-2-yl}amino)-2-methylbenzenesulfonamide



Procedure 1

To a solution of Intermediate Example 4 (200 mg, 0.695 mmol) and 5-amino-2-methylbenzenesulfonamide (129.4 mg, 0.695 mmol) in isopropanol (6 ml) was added 4 drops of conc. HCl. The mixture was heated to reflux overnight. The mixture was cooled to rt and diluted with ether (6 ml). Precipitate was collected via filtration and washed with ether. The hydrochloride salt of 5-({4-[(2,3-dimethyl-2*H*-indazol-6-yl)(methyl)amino]-pyrimidin-2-yl}amino)-2-methylbenzenesulfonamide was

isolated as an off-white solid. ¹H NMR (400 MHz, d₆DMSO+NaHCO₃) δ 9.50 (br s, 1H), 8.55 (br s, 1H), 7.81 (d, *J* = 6.2 Hz, 1H), 7.75 (d, *J* = 8.7 Hz, 1H), 7.69 (m, 1H), 7.43 (s, 1H), 7.23 (s, 2H), 7.15 (d, *J* = 8.4 Hz, 1H), 6.86 (m, 1H), 5.74 (d, *J* = 6.1 Hz, 1H), 4.04 (s, 3H), 3.48 (s, 3H), 2.61 (s, 3H), 2.48 (s, 3H). MS (ES+, *m/z*) 438 (M+H).

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Procedure 2

A 250-mL 3-necked flask equipped with a magnetic stir bar, thermometer, reflux condenser, and nitrogen inlet/outlet was charged with ethanol (60 mL, 10 volumes), the product of Intermediate Example 4 (6.00 g, 20.85 mmol, 1.0 equiv) and 5-amino-2-methylbenzenesulfonamide (4.00 g, 21.48 mmol, 1.03 equiv) with stirring. The reaction mixture was heated to 70 °C. After stirring the reaction mixture at 68 – 72 °C for 3 hrs, 4M HCl in dioxane (0.11 mL, 0.44 mmol, 0.02 equiv) was charged over ca. 2 min. The reaction mixture was stirred at 68 - 72 °C until < 1.5% by area of the starting product of Intermediate Example 4 was remaining by HPLC analysis (Typically, this reaction is complete in > 8 hrs). The reaction mixture was cooled to 20 °C over ca. 30 min and stirred at 20 - 22 °C for 40 min. The product was then isolated by filtration and the filter cake washed with ethanol (20 mL, 3.3 volumes). The wet cake was dried under vacuum at 45 - 50 °C. The monohydrochloride salt of 5-({4-[(2,3-dimethyl-2*H*-indazol-6-yl)(methyl)amino]-pyrimidin-2-yl}amino)-2-methylbenzenesulfonamide (9.52 g, 96.4%) was isolated as a white solid. ¹H NMR (400 MHz, d₆DMSO+NaHCO₃) δ 9.50 (br s, 1H), 8.55 (br s, 1H), 7.81 (d, *J* = 6.2 Hz, 1H), 7.75 (d, *J* = 8.7 Hz, 1H), 7.69 (m, 1H), 7.43 (s, 1H), 7.23 (s, 2H), 7.15 (d, *J* = 8.4 Hz, 1H), 6.86 (m, 1H), 5.74 (d, *J* = 6.1 Hz, 1H), 4.04 (s, 3H), 3.48 (s, 3H), 2.61 (s, 3H), 2.48 (s, 3H). MS (ES+, *m/z*) 438 (M+H).

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Procedure 3:

To a stirred suspension of the product of Intermediate Example 4 (1.1 g, 3.8 mmol) in 14 mL of MeOH, was added 5-amino-2-methylbenzenesulfonamide (0.78 g, 4.2 mmol, 1.1 equiv) at room temperature. The reaction mixture was heated at reflux for 3 h, then 4 M HCl in 1,4-dioxane (19 μL, 0.076 mmol) was added in one portion. After 4 h, the suspension was cooled to room temperature, and filtered. The resulting solid was washed with 10 mL of MeOH and dried in vacuo to yield 1.3 g (72%) of 5-({4-[(2,3-dimethyl-2*H*-indazol-6-yl)methylamino]-2-pyrimidinyl}amino)-2-methyl benzenesulfonamide monohydrochloride as a white solid. ¹H NMR (DMSO-d₆, 400 MHz) δ 10.95 (s, 1H), 8.36 (s, 1H), 7.86 (d, *J* = 8.8 Hz, 2H), 7.64-7.59 (m,

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2H), 7.40 (m, 3H), 6.93 (dd, $J = 8.8, 2.0$ Hz, 1H), 5.92 (s, 1H), 4.08 (s, 3H), 3.57 (s, 3H), 2.65 (s, 3H), 2.56 (s, 3H).

Procedure 4

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To a stirred suspension of the product of Intermediate Example 4 (1.1 g, 3.7 mmol) in 10 mL of THF, was added 5-amino-2-methylbenzenesulfonamide (0.70 g, 3.8 mmol, 1.0 equiv) at room temperature. The reaction mixture was heated at reflux for 3 h, then 4 M HCl in 1,4-dioxane (18 μ L, 0.072 mmol) was added in one portion.

10 After 5 h, the suspension was cooled to room temperature, and filtered. The resulting solid was washed with 16 mL of THF and dried in the air to yield 1.6 g (92%) of 5-({4-[(2,3-dimethyl-2H-indazol-6-yl)methylamino]-2-pyrimidinyl}amino)-2-methylbenzene sulfonamide monohydrochloride as a light yellow solid.

15 Procedure 5

To a stirred suspension of the product of Intermediate Example 4 (1.0 g, 3.6 mmol) in 10 mL of CH₃CN, was added 5-amino-2-methylbenzenesulfonamide (0.70 g, 3.8 mmol, 1.0equiv) at room temperature. The reaction mixture was heated at reflux for 3 h, then 4 M HCl in 1,4-dioxane (18 μ L, 0.076 mmol) was added in one portion. After 20 h, the suspension was cooled to room temperature, and filtered. The resulting solid was washed with 10 mL of CH₃CN and dried in the air to yield 1.3 g (73%) of 5-({4-[(2,3-dimethyl-2H-indazol-6-yl)methylamino]-2-pyrimidinyl}amino)-2-methyl benzenesulfonamide monohydrochloride as an off-white solid.

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Procedure 6

Preparation of 5-({4-[(2,3-dimethyl-2H-indazol-6-yl) (methyl) amino] pyrimidin-2-yl}amino)-2-methylbenzenesulfonamide methanesulfonic acid salt.

30 In a 250 mL flask the product of Example 1, procedure 1, (1.0 g, 2.29 mmol) was slurried in water (19 mL). Methanesulfonic acid (0.231 g, 2.4 mmol) was added all at once and the mixture was heated to reflux for 5 min. The mixture was cooled to 0 °C over a 1 hour period and was then isolated by filtration and air dried. 5-({4-[(2,3-Dimethyl-2H-indazol-6-yl) (methyl) amino] pyrimidin-2-yl}amino)-2-methylbenzenesulfonamide methanesulfonic acid salt (1.03 g, 84%) was obtained as
35 a white solid. mp = 247-248 °C.

Procedure 7:

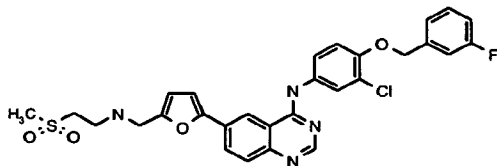
Preparation of 5-({4-[(2,3-dimethyl-2H-indazol-6-yl)methylamino]-2-pyrimidinyl}amino)-2-methylbenzenesulfonamide monohydrochloride monohydrate.

5 To a round bottom flask, was added 2.6 g of the monohydrochloride salt of Example 1, procedure 1, any form. Then added was 39 mL of isopropanol (15 volumes). The mixture was heated to 75 deg C in an oil bath, then 14 mL of 0.05N aqueous HCl (5.4 volumes) was added. The clear solution was cooled to 65 deg C, then seeded with the monohydrate of the monohydrochloride salt of Example 1,
10 procedure 1 (0.05-0.1 wt %). The cloudy solution was stirred at 65 deg C for 60 minutes, then cooled to 0 deg C at ~0.25-0.5 deg C/min. The resulting white solid was filtered and dried to constant weight under vacuum at RT to give 88% yield of 5 - ({4-[(2,3-dimethyl-2H-indazol-6-yl)methylamino]-2-pyrimidinyl}amino)-2-methylbenzene sulfonamide monohydrochloride monohydrate.

Example 2

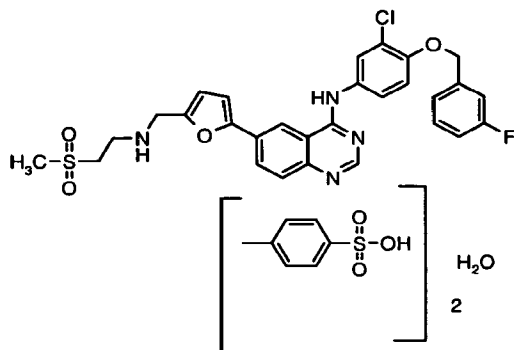
Monohydrate ditosylate salt of N-{3-Chloro-4-[(3-fluorobenzyl)oxy]phenyl}-6-[5-({[2-(methane sulphonyl) ethyl]amino}methyl)-2-furyl]-4-quinazolinamine (monohydrate ditosylate salt of compound of formula (II))

1(a) Preparation of N-{3-Chloro-4-[(3-fluorobenzyl)oxy]phenyl}-6-[5-({[2-(methane sulphonyl) ethyl]amino}methyl)-2-furyl]-4-quinazolinamine (free base of compound of formula (II))



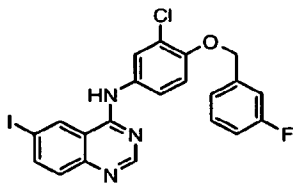
The title compound was prepared according to Procedure D of International Applications WO 02/02552: p. 16, line 19 to p. 17, line 3 and WO 99/35146: p. 56,
30 lines 20-32 and Example 29 p. 100, lines 18-29, from 5-(4-{3-chloro-4-(3-fluorobenzyloxy)-anilino}-6-quinazolinyl)-furan-2-carbaldehyde (0.6 equiv) and 2-methanesulphonyl-ethylamine (1 equiv). ¹H NMR 400 MHz (DMSO-d₆) 9.60 (bs, 1H); 9.32 (bs, 1H); 8.82 (bs, 1H); 8.34 (d, 1H); 8.0 (s, 1H); 7.88 (d, 1H); 7.74 (d, 1H); 7.45 (m, 1H); 7.34-7.23 (m, 4H); 7.17 (m, 1H); 6.83 (d, 1H); 5.27 (s, 2H); 4.42 (s, 2H);
35 3.59 (m, 2H); 3.40 (m, 2H, obscured by waterpeak); 3.12 (s, 3H); MS *m/z* 581 (M+H⁺).

1(b) Preparation of monohydrate ditosylate salt of N-{3-Chloro-4-[(3-fluorobenzyl)oxy]phenyl}-6-[5-({[2-(methane sulphonyl) ethyl]amino)methyl]-2-furyl]-4-quinazolinamine (monohydrate ditosylate salt of compound of formula (II))



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Stage 1: Preparation of N-{3-chloro-4-[(3-fluorobenzyl)oxy]phenyl}-6-iodo-4-quinazolinamine

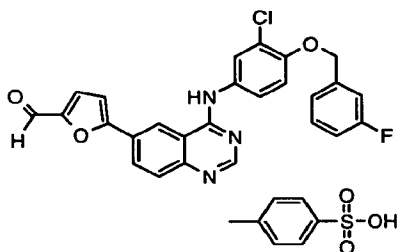


- 4-Chloro-6-iodoquinazoline (1wt) was added to a solution of fluorobenzylloxylaniline (0.894wt, 1.03equiv) in N-methylpyrrolidinone (8.26wt, 8vol) at *ca* 20°C, and after the initial exotherm had subsided, the resulting solution was stirred at 20°-25°C for at least 30 minutes. The dark solution was treated with triethylamine (0.58vol, 1.2equiv) and the mixture was stirred for 20-30 minutes.
- Isopropanol (2.5vol) was added and the mixture was heated to *ca* 50°C. Water (up to 3vol) was added slowly to the vessel over 10-15 minutes, while keeping the temperature at *ca* 50°C. Once crystallisation had commenced the addition was stopped and the resulting slurry was aged for 30-45 minutes at *ca* 50°C. Any residual water (from the 3vol) was added, then further water (5vol) was added to the vessel over *ca* 30 minutes while maintaining the temperature at *ca* 50°C. The resulting slurry was cooled to *ca* 20°C over *ca* 30 minutes and aged at *ca* 20°C for at least 30 minutes. The solid was collected by filtration and washed sequentially with

water (2 x 5vol), then isopropanol (5vol). The product was dried *in vacuo* at *ca* 60°C to give the title compound as a cream crystalline solid.

Stage 2: Preparation of 5-(4-[3-chloro-4-(3-fluorobenzoyloxy)-anilino]-6-quinazoliny)-furan-2-carbaldehyde 4-methylbenzenesulfonate

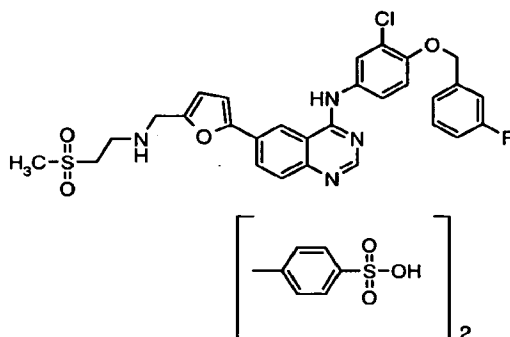
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A mixture of N-{3-chloro-4-[(3-fluorobenzyl)oxy]phenyl}-6-iodo-4-quinazolinamine (1wt), boronic acid (0.37wt, 1.35equiv), and 10% palladium on charcoal (0.028wt, 50% water wet) was slurried in IMS (15vol). The resultant suspension was stirred for 5 minutes, treated with di-isopropylethylamine (0.39vol, 1.15equiv) and then heated to *ca* 70°C for *ca* 3 hours when the reaction was complete (determined by HPLC analysis). The mixture was diluted with tetrahydrofuran (THF, 15vol) and then filtered (hot - through GFA filter paper) to remove catalyst. The vessel was rinsed with IMS (2vol).

A solution of p-toluenesulfonic acid monohydrate (1.54wt, 4.1equiv) in water (3vol) was added over 5-10 minutes to the filtered solution maintained at 65°C. After crystallisation the suspension was stirred at 60°-65°C for 1 hour, cooled to *ca* 25°C over 1 hour and stirred at this temperature for a further 2 hours. The solid was collected by filtration, washed with IMS (3vol) then dried *in vacuo* at *ca* 50°C to give the title compound as a yellow-orange crystalline solid.

Stage 3: Preparation of anhydrous ditosylate salt of N-{3-Chloro-4-[(3-fluorobenzyl)oxy]phenyl}-6-[5-({[2-(methane sulphonyl) ethyl]amino)methyl}-2-furyl]-4-quinazolinamine (anhydrous ditosylate salt of compound of formula (II))

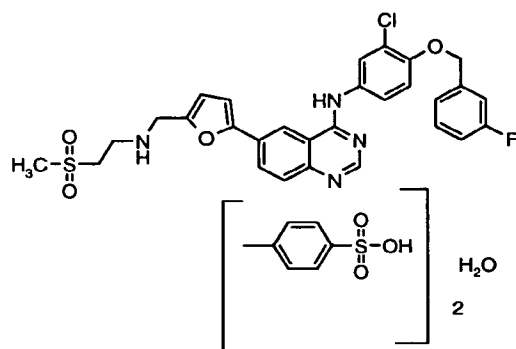


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5-(4-[3-chloro-4-(3-fluorobenzyloxy)-anilino]-6-quinazolinyl)-furan-2-carbaldehyde 4-methylbenzenesulfonate (1 wt) and 2-(methylsulfonyl) ethylamine hydrochloride (0.4 wt, 1.6equiv) were suspended in THF (10vol). Sequentially, acetic acid (0.35vol, 4equiv) and di-isopropylethylamine (1.08vol, 4equiv) were added. The resulting solution was stirred at 30°-35°C for ca 1 hour then cooled to ca 23°C. Sodium triacetoxymethylborohydride (0.66wt, 2equiv) was then added as a continual charge over approximately 15 minutes (some effervescence is seen at this point). The resulting mixture was stirred at ca 22°C for ca 2 hours then sampled for HPLC analysis. The reaction was quenched by addition of 5M aqueous sodium hydroxide (5vol) and stirred for ca 30 minutes (some effervescence is seen at the start of the caustic addition).

The aqueous phase was then separated, extracted with THF (2vol) and the combined THF extracts were then washed with 10%w/v aqueous sodium chloride solution (4vol). A solution of *p*-toluenesulfonic acid monohydrate (pTSA, 1.77wt, 6equiv) in THF (7 vol)¹ was prepared and warmed to ca 55°C. The THF solution of N-{3-Chloro-4-[(3-fluorobenzyl)oxy]phenyl}-6-[5-({[2-(methanesulphonyl) ethyl] amino)methyl}- 2-furyl]-4-quinazolinamine was added to the pTSA solution over at least 30minutes, maintaining the batch temperature at ca 55±3°C². The resulting suspension was stirred at ca 55°C for 2 hours, cooled to 20°-25°C over ca 60 minutes and aged at this temperature for ca 30 minutes. The solid was collected by filtration, washed with THF (2 x 2vol) and dried *in vacuo* at ca 40°C to give the desired compound as a pale yellow crystalline solid.

Stage 4: Preparation of monohydrate ditosylate salt of N-{3-Chloro-4-[(3-fluorobenzyl)oxy]phenyl}-6-[5-({[2-(methane sulphonyl) ethyl]amino)methyl}-2-furyl]-4-quinazolinamine (monohydrate ditosylate salt of compound of formula (II))



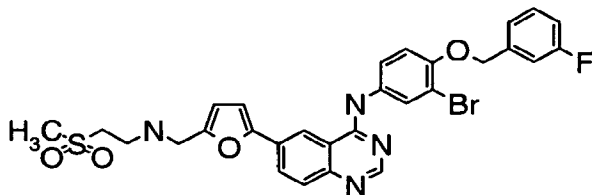
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A suspension of the anhydrous ditosylate salt of N-{3-Chloro-4-[(3-fluorobenzyl)oxy]phenyl}-6-[5-({[2-(methane sulphonyl) ethyl]amino)methyl}-2-furyl]-4-quinazolinamine (1 wt), in tetrahydrofuran (THF, 14 vol) and water (6 vol) was heated to *ca* 55°-60°C for 30 minutes to give a solution which was clarified by
 10 filtration and the lines washed into the crystallisation vessel with THF/Water (7:3, 2 vol). The resultant solution was heated to reflux and tetrahydrofuran (9 vol, 95% w/w azeotrope with water) was distilled off at atmospheric pressure.

The solution was seeded with N-{3-Chloro-4-[(3-fluorobenzyl)oxy]phenyl}-6-[5-({[2-(methane sulphonyl) ethyl]amino)methyl}-2-furyl]-4-quinazolinamine ditosylate
 15 monohydrate (0.002 wt). Once the crystallisation was established water (6 vol) was added while maintaining the reaction temperature above 55°C. The mixture was cooled to 5°-15°C over *ca* 2 hours. The solid was collected by filtration, washed with tetrahydrofuran/water (3:7 ratio, 2 x 2 vol) and dried *in vacuo* at 45°C to give N-{3-Chloro-4-[(3-fluorobenzyl)oxy]phenyl}-6-[5-({[2-(methane sulphonyl) ethyl]
 20 amino)methyl}-2-furyl]-4-quinazolinamine ditosylate monohydrate as a bright yellow crystalline solid.

Example 3

Preparation of (4-(3-Fluoro-benzyloxy-3-bromophenyl)-(6-(5-((2-methanesulphonyl-ethylamino)-methyl)-furan-2-yl)-quinazolin-4-yl)-amine and di-hydrochloride and ditosylate salts thereof

*(a) Preparation of 2-Bromo-4-nitrophenol*

2-Bromo-4-nitroanisole (20 g, 0.086 mol) was dissolved in DMF (414 mL) at room temperature under N₂. Sodium ethylthiolate (17.4 g, 0.207 mol) was added and the reaction mixture was warmed to 115°C for 2 hours. The reaction was cooled to room temperature and diluted with EtOAc (200 mL) and 1 M HCl (aq., 200 mL). The phases were separated, and the desired product was extracted into 1 M NaOH (aq, 150 mL X 3). The basic aqueous extracts were combined and acidified using conc. HCl. The desired product was extracted from the acidic aqueous solution using EtOAc (250 mL X 2). The combined organic layers were washed with brine and dried over sodium sulfate. The volatiles were removed in vacuo to afford a light brown semi-solid (9.8 g, 52% yield). ¹H NMR (DMSO-d₆) δ 8.33 (m, 1H); 8.09 (m, 1H); 7.07 (d, 1H).

(b) Preparation of 2-Bromo-1-(3-fluorobenzyloxy)-4-nitrobenzene

2-Bromo-4-nitrophenol (4.86 g, 0.0223 mol), triphenylphosphine (7.6 g, 0.0290 mol), 3-fluorobenzylalcohol (3.65 g, 0.0290 mol) were combined and dissolved in THF (89 mL). The reaction temperature was cooled to 0°C and DIAD (4.50 g, 0.0290 mol) was added. The reaction was allowed to warm slowly to room temperature and stirred for 3 hours before it was diluted with water (100 mL) and EtOAc (100 mL). The layers were separated and the aqueous layer was extracted with EtOAc (200 mL X 2). The organic extracts were combined and washed with brine, followed by drying over sodium sulfate. The volatiles were removed in vacuo and the residual semi-solid was treated with diethyl ether. The solids were removed

by filtration. The volatiles from the resulting filtrate were removed in vacuo and the material was purified using EtOAc:Hexanes (90/10) in a biotage LC system to afford the title compound as a yellow solid (3.73 g, 68% yield). ¹H NMR (DMSO-d₆) δ 8.43 (d, 1H); 8.26 (m, 1H); 7.45 (m, 1H); 7.38 (d, 1H); 7.30 (m, 2H); 7.17 (m, 1H); 5.39 (s, 2H).

(c) Preparation of 3-Bromo-4-(3-fluorobenzyloxy)-aniline

Under a blanket of N₂, Pt/C (5%, 0.37 g) was charged to a Parr Shaker Flask. Ethanol (150 mL) and 2-bromo-1-(3-fluorobenzyloxy)-4-nitrobenzene (3.73 g, 0.011 mol) were added and the reaction mixture was placed on a Parr Shaker Apparatus under 30 psi of H₂ for 5 h. The reaction was filtered through a pad of Celite to remove the catalyst and the volatiles were removed from the filtrate. The residue was dissolved in the CH₂Cl₂ (5 mL) and treated with conc. HCl (1 mL). The precipitate was collected by filtration and free-based using saturated aqueous sodium bicarbonate (2.27g, 67% yield) ¹H NMR (DMSO-d₆) δ 7.4 (m, 1H); 7.23 (m, 2H); 7.11 (m, 1H); 6.86 (d, 1H); 6.77 (m, 1H); 6.48 (m, 1H); 5.0 (s, 2H); 4.93 (bs, 2H).

(d) Preparation of 6-Iodo-(4-(3-fluorobenzyloxy)-3-bromophenyl)-quinazolin-4-yl)amine

The title compound was prepared according to Procedure A from 3-bromo-4-(3-fluorobenzyloxy)-aniline (0.79 g, 2.7 mmol) and 4-chloro-6-iodo-quinazoline (0.8g, 2.7 mmol). ¹H NMR (DMSO-d₆) δ 11.1 (bs, 1H); 9.10 (s, 1H); 8.87 (s, 1H); 8.29 (d, 1H); 8.03 (s, 1H); 7.68 (m, 1H); 7.62 (d, 1H); 7.45 (m, 1H); 7.33-7.26 (m, 3H); 7.16 (m, 1H); 5.28 (s, 2H).

(e) Preparation 5-(4-(3-Bromo-4-(3-fluorobenzyloxy)-anilino)-quinazolin-6-yl)-furan-2-carbaldehyde

The title compound was prepared according to Procedure B followed by Procedure C from 6-iodo-(4-(3-fluorobenzyloxy)-3-bromophenyl)-quinazolin-4-yl)amine (1.0 g, 1.82 mmol) and (1,3 dioxolan-2-yl)-2-(tributylstannyl)furan (1.17 g, 2.73 mmol). ¹H NMR (DMSO-d₆) δ 11.89 (bs, 1H); 9.66 (s, 1H); 9.41 (s, 1H); 8.90 (s, 1H); 8.49 (d, 1H); 8.05 (m, 1H); 7.96 (d, 1H); 7.75 (m, 1H); 7.70 (m, 1H); 7.61 (m, 1H); 7.43 (m, 1H); 7.30 (m, 3H); 7.16 (m, 1H); 5.29 (s, 2H).

(f) Preparation of (4-(3-Fluoro-benzyloxy-3-bromophenyl)-(6-(5-((2-methanesulphonyl-ethylamino)-methyl)-furan-2-yl)-quinazolin-4-yl)-amine dihydrochloride

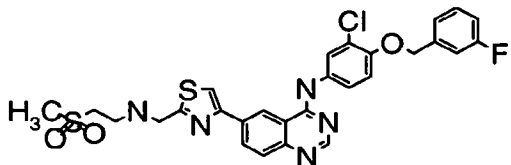
5 The title compound was prepared according to Procedure D from a mixture of 5-(4-(3-Bromo-4-(3-fluorobenzyloxy)-anilino)-quinazolin-6-yl)-furan-2-carbaldehyde (0.623 g, 1.2 mmol) in dichloroethane (12 mL), triethylamine (0.167 mL, 1.2 mmol), acetic acid (0.216 mL 3.6 mmol), and 2-methanesulphonylethylamine (0.447 g, 3.6 mmol). The reaction mixture was warmed to reflux for 1 hour and then cooled to rt before adding sodium triacetoxyborohydride (0.5 g). After 0.5 hours of stirring, another aliquot of sodium triacetoxyborohydride (0.5 g) was added and the reaction was stirred an additional 0.5 hours. The reaction was quenched by the addition of a saturated solution of sodium bicarbonate (aq, 50 mL). EtOAc (50 mL) was added and the layers were separated. The organics were washed with brine and dried over sodium sulfate. The volatiles were removed in vacuo. Purification of the compound was achieved using Biotage column chromatography; eluents: CH₂Cl₂, EtOH, Et₃N (150:8:1). The appropriate fractions were combined and the volatiles were removed in vacuo. The compound was crystallized from EtOAc and Et₂O to afford a yellow solid. The hydrochloride salt was made by dissolving the material in a minimal amount of EtOAc and adding 2M HCl in diethyl ether (0.5 mL) to afford a dark yellow solid (0.27 g, 35% yield). ¹H NMR (DMSO-d₆) δ 11.70 (bs, 1H); 9.84 (bs, 2H); 9.59 (s, 1H); 8.89 (s, 1H); 8.39 (d, 1H); 8.14 (s, 1H); 7.93 (d, 1H); 7.80 (d, 1H); 7.45 (m, 1H); 7.31 (m, 4H); 7.16 (m, 1H); 6.83 (m, 1H); 5.30 (s, 2H); 4.43 (s, 2H); 3.67 (m, 2H); 3.40 (m, 2H); 3.12 (s, 3H).

(g) Preparation of (4-(3-Fluoro-benzyloxy)-3-bromophenyl)-(6-(5-((2-methanesulphonyl-ethylamino)-methyl)-furan-2-yl)-quinazolin-4-yl)-amine ditosylate.

30 The HCl salt of 5-(4-[3-bromo-4-(3-fluorobenzyloxy)-anilino]-6-quinazolinyl)-furan-2-carbaldehyde, is prepared according to Procedure D and Example 1(e), and is converted to the tosylate salt according to the procedure of Example 1(h). The resultant carbaldehyde tosylate product is used to prepare the (4-(3-Fluoro-benzyloxy)-3-bromophenyl)-(6-(5-((2-methanesulphonyl-ethylamino)-methyl)-furan-2-yl)-quinazolin-4-yl)-amine ditosylate according to the procedure of Example 1(i).

Example 4

Preparation of (4-(3-Fluoro-benzyloxy)-3-chlorophenyl)-(6-(2-((2-methanesulphonyl-ethylamino)-methyl)-thiazol-4-yl)quinazolin-4-yl)-amine and di-hydrochloride and ditosylate salts thereof



(a) Preparation of N-(4-(3-fluorobenzyloxy)-chlorophenyl)-6-(1-ethoxyvinylether)-quinazolin-4-yl)-amine

To a suspension of the 6-iodo-(4-(3-fluorobenzyloxy)-3-chlorophenyl)-quinazolin-4-yl amine (12.6 g, 24.93 mmol) in acetonitrile (100 mL) was added tributyl(1-ethoxyvinyl)stannane (9 g, 24.93 mmol) and bis(triphenylphosphine) palladium (II) chloride (1.75 g, 2.29 mmol). The reaction mixture was refluxed for 18 hours, then filtered through a plug of silica gel. The resulting solution was poured into 5% aqueous NH_4OH (200 mL) and extracted with ethyl acetate (500 mL). The organic layer was dried over anhydrous sodium sulfate, concentrated, and purified by silica gel chromatography to provide the title compound as a yellow solid (7.2 g, 64% yield). ^1H NMR (400 MHz, d_6 DMSO) δ 9.92 (s, 1H), 8.76 (s, 1H), 8.58 (s, 1H), 8.08 (m, 1H), 8.01 (m, 1H), 7.76 (m, 2H), 7.48 (m, 1H), 7.32 (m, 3H), 7.22 (m, 1H), 5.28 (s, 2H), 5.02 (s, 1H), 4.56 (s, 1H), 4.01 (q, 2H), 1.42 (t, 3H); ESI-MS m/z 449.9 ($\text{M}+\text{H}$) $^+$.

(b) Preparation of N-{4-[(3-fluorobenzyloxy)]-chlorophenyl}-6-[2-[[2-(methanesulphonyl)ethyl]-[trifluoroacetyl]amino]methyl]-1,3-thiazol-4-yl]-quinazolin-4-yl)-amine

To a solution of N-(4-(3-fluorobenzyloxy)-chlorophenyl)-6-(1-ethoxyvinylether)-quinazolin-4-yl)-amine (7.1 g, 15.8 mmol) in a THF (150 mL)/ H_2O (5 mL) mixture cooled to 0°C was added N-bromosuccinimide (2.81 g, 15.8 mmol). The resulting mixture was stirred for 0.25 hours, then dried over anhydrous sodium sulfate and concentrated. The crude N-(4-(3-fluorobenzyloxy)-chlorophenyl)-6-(bromomethylketone)-quinazolin-4-yl)-amine and N-(trifluoroacetyl)-N-(methanesulphonylethyl)-aminomethylthioamide (4.61g, 15.8 mmol) were dissolved in DMF (50 mL) and heated at 70°C for 1 hour. The reaction mixture was

concentrated, then diluted with dichloromethane (300 mL) and washed with saturated sodium bicarbonate solution (100 mL). The organic layer was dried over anhydrous sodium sulfate, concentrated, and purified by silica gel chromatography to provide the title compound as a foam (4.6 g, 42% yield). ESI-MS m/z 694.1 (M+H)⁺.

5

(c) Preparation of N-{4-[(3-fluorobenzyloxy)]-chlorophenyl}-6-[2-((2-methanesulphonyl)ethyl)-amino)methyl]-1,3-thiazol-4-yl]-quinazolin-4-yl)-amine hydrochloride

10 To a solution of N-{4-[(3-fluorobenzyloxy)]-chlorophenyl}-6-[2-((2-methanesulphonyl)ethyl)-[trifluoroacetyl]amino)methyl]-1,3-thiazol-4-yl]-quinazolin-4-yl)-amine (4.6 g, 6.63 mmol) in methanol (100 mL) was added 2M NaOH (50 mL). The resulting mixture was stirred at room temperature for 2 hours, concentrated to ½ volume, poured into H₂O (100 mL), and extracted with dichloromethane (300 mL).
 15 The organic layer was dried over anhydrous sodium sulfate, concentrated, and purified by silica gel chromatography. The resulting amine was dissolved in dichloromethane/methanol (3:1, 100 mL) and then 4M HCl/dioxane (20 mL) was added. The resulting mixture was concentrated and filtered to provide the title compound as a yellow solid (4.0 g, 90% yield). ¹H NMR (400 MHz, d₄ MeOH) δ 9.38
 20 (s, 1H), 8.82 (s, 1H), 8.78 (d, 1H), 8.36 (s, 1H), 7.94 (s, 1H), 7.92 (d, 1H), 7.63 (m, 1H), 7.41 (m, 1H), 7.26 (m, 1H), 7.22 (m, 2H), 7.04 (m, 1H), 5.24 (s, 2H), 4.82 (s, 2H), 3.84 (m, 2H), 3.76 (m, 2H), 3.12 (s, 3H); ESI-MS m/z 597.1 (M+H)⁺.

25 *(d) Preparation of (4-(3-Fluoro-benzyloxy)-3-chlorophenyl)-(6-(2-((2-methanesulphonyl-ethylamino)-methyl)-thiazol-4-yl)quinazolin-4-yl)-amine ditosylate.*

The HCL salt of (4-(3-Fluoro-benzyloxy)-3-chlorophenyl)-(6-(2-((2-methanesulphonyl-ethylamino)-methyl)-thiazol-4-yl)quinazolin-4-yl)-amine was prepared according to Procedures 3(a) to (c) and then converted to the (4-(3-Fluoro-benzyloxy)-3-chlorophenyl)-(6-(2-((2-methanesulphonyl-ethylamino)-methyl)-thiazol-4-yl)quinazolin-4-yl)-amine ditosylate salt according to the procedure of Examples 1 and 2. ¹H NMR (300 MHz, d₆-DMSO) 11.4 (br s, 1H), 9.51 (br s, 1H), 9.24 (s, 1H), 8.95 (s, 1H), 8.68 (d, J = 9 Hz, 1H), 8.42 (s, 1H), 7.96 (d, J = 9 Hz, 1H), 7.89 (d, J = 2 Hz, 1H), 7.64 (dd, J = 2, 9 Hz, 1H), 7.47 (m, 5H), 7.34 (m, 3H), 7.20 (t, J = 9Hz, 1H), 7.10 (d, J = 8Hz, 4H), 5.32 (s, 2H), 4.76 (d, 2H), 3.61 (s, 4H), 3.15 (s, 3H), 2.28 (s, 6H).

35

Biological Data**Tumor Studies: BT474**

The BT474 xenografts were maintained by serial transplantation of tumor
5 fragments in SCID mice. Tumors are initiated by injection of tumor fragments
subcutaneously in the axillary region.

Tumor Studies: NCI H322

NCI H322 cells obtained from the ATCC and were cultured in RPMI 1640 +
10 10% Fetal bovine serum, Sodium pyruvate and L-Glutamine at 37° in a 95/5%
air/CO₂ atmosphere. Cells were harvested following trypsin digestion and brought to
a density of 2x10⁶ cells / 200 µl in PBS. Tumors were initiated by injection of the cell
suspension subcutaneously in the axillary region. In addition some experiments were
performed following serial transplantation of tumor fragments in SCID mice. Tumors
15 were initiated by injection of tumor fragments subcutaneously in the axillary region.

Tumor Studies: Measurements

For the xenograft models used here, solid tumors were measured by
electronic caliper measurement through the skin. Measurements were typically made
20 twice weekly.

Tumor Studies: Formulation and Administration

Drugs were administered by P.O. route. The compounds or salts thereof of
Example 1 and Example 2 were formulated in aqueous 0.5% hydroxypropyl
25 methylcellulose, 0.1% Tween 80 and administered as a suspension once daily for 21
days as indicated in the respective figures. These studies were performed under
IACUC # 468 and 603. The results are illustrated in Figures 1- 2.

Figure 1 is a representative experiment and illustrates dosing of a BT474
30 (breast) subcutaneous human xenograft mouse model with the compound of
Example 1 and/or the compound of Example 2. The compound of Example 1 as a
monotherapy in the BT474 s.c. human xenograft mouse model showed some anti-
tumor activity (about 16-78% tumor growth inhibition). Dosing of the compound of
Example 2 also showed anti-tumor activity in the same model as monotherapy
35 (about 21-99% tumor growth inhibition at highest dose). When the compound of

Example 1 and the compound of Example 2 were used in combination , 79-109% tumor growth inhibition was observed during treatment.

5 Table 1 summarizes the results of two independent experiments of dosing of a BT474 (breast) subcutaneous human xenograft mouse model with the compound of Example 1 and/or the compound of Example 2. The column labeled "Exp.1" contains the results presented in Figure 1 for the last data point on the graph. The column labeled "Exp. 2" are analogous results from an independent experiment.

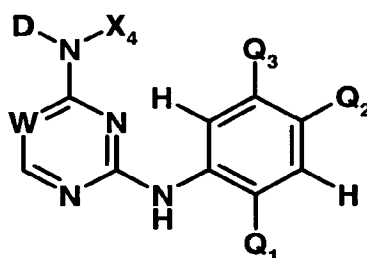
10 Figure 2 illustrates dosing of a NCI H322 (non-small cell lung carcinoma) subcutaneous human xenograft mouse model with the compound of Example 1 and/or the compound of Example 2. The compound of Example 1 dosed as monotherapy in the NCI H322 s.c. human xenograft mouse model showed some anti-tumor activity (about 86-88% tumor growth inhibition). The compound of
15 Example 2 also showed anti-tumor activity in the same model when dosed as monotherapy (about 28-54% tumor growth inhibition at the highest dose tested). When the compound of Example 1 and the compound of Example 2 were used in combination, 60-108% anti-tumor activity was observed during treatment.

20 Table 2 summarizes the results of three independent experiments of dosing of a NCI H322 (non-small cell lung carcinoma) subcutaneous human xenograft mouse model with the compound of Example 1 and/or the compound of Example 2. The column labeled "Exp.3" contains the results presented in Figure 2 for the last data point on the graph. The columns labeled "Exp. 1" and "Exp. 2" are analogous
25 results from independent experiments.

CLAIMS

We claim:

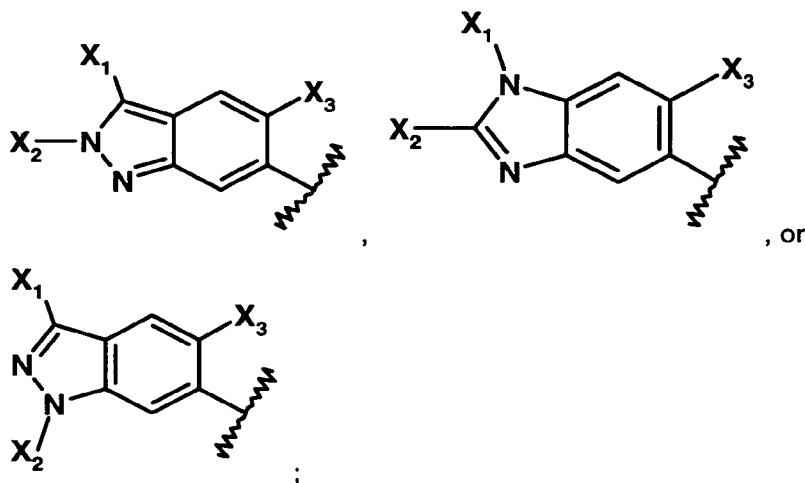
1. A method of treating cancer in a mammal, comprising: administering to said
 5 mammal
 (a) a compound of formula I,



(I)

- 10 or a salt, solvate, or physiologically functional derivative thereof;
 wherein:

D is



15

X₁ is hydrogen, C₁₋₄ alkyl, C₁₋₄ haloalkyl, or C₁₋₄ hydroxyalkyl;

X₂ is hydrogen, C₁₋₄ alkyl, C₁₋₄ haloalkyl, C(O)R¹, or aralkyl;

X₃ is hydrogen or halogen;

- 20 X₄ is hydrogen, C₁₋₄ alkyl, C₁₋₄ haloalkyl, heteroaralkyl, cyanoalkyl,
 -(CH₂)_pC=CH(CH₂)_iH, -(CH₂)_pC≡C(CH₂)_iH, or C₃₋₇ cycloalkyl;

p is 1, 2, or 3;

t is 0 or 1;

W is N or C-R, wherein R is hydrogen, halogen, or cyano;

Q₁ is hydrogen, halogen, C₁₋₂ haloalkyl, C₁₋₂ alkyl, C₁₋₂ alkoxy, or C₁₋₂ haloalkoxy;

5 Q₂ is A¹ or A²;

Q₃ is A¹ when Q₂ is A² and Q₃ is A² when Q₂ is A¹;

wherein

A¹ is hydrogen, halogen, C₁₋₃ alkyl, C₁₋₃ haloalkyl, -OR¹, and

A² is the group defined by -(Z)_m-(Z¹)-(Z²), wherein

10 Z is CH₂ and m is 0, 1, 2, or 3, or

Z is NR² and m is 0 or 1, or

Z is oxygen and m is 0 or 1, or

Z is CH₂NR² and m is 0 or 1;

Z¹ is S(O)₂, S(O), or C(O); and

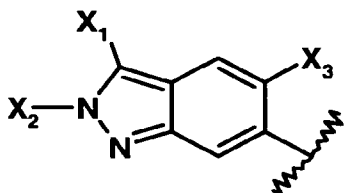
15 Z² is C₁₋₄ alkyl, NR³R⁴, aryl, arylamino, aralkyl, aralkoxy, or heteroaryl,

R¹ is C₁₋₄ alkyl;

R², R³, and R⁴ are each independently selected from hydrogen, C₁₋₄ alkyl, C₃₋₇ cycloalkyl, -S(O)₂R⁵, and -C(O)R⁵;

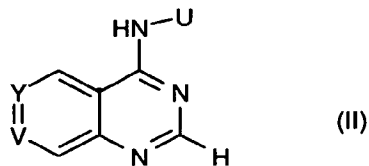
20 R⁵ is C₁₋₄ alkyl, or C₃₋₇ cycloalkyl; and

when Z is oxygen then Z¹ is S(O)₂ and when D is



then X₂ is C₁₋₄ alkyl, C₁₋₄ haloalkyl, C(O)R¹, or aralkyl; and

(b) a compound of formula II:



25

or a salt, solvate, or physiologically functional derivative thereof;

wherein

Y is CR¹ and V is N;

or Y is CR¹ and V is CR²;

- R¹ represents a group CH₃SO₂CH₂CH₂NHCH₂-Ar-, wherein Ar is selected from phenyl, furan, thiophene, pyrrole and thiazole, each of which may optionally be substituted by one or two halo, C₁₋₄ alkyl or C₁₋₄ alkoxy groups;

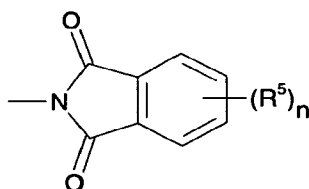
R² is selected from the group comprising hydrogen, halo, hydroxy, C₁₋₄ alkyl, C₁₋₄ alkoxy, C₁₋₄ alkylamino and di[C₁₋₄ alkyl]amino;

- U represents a phenyl, pyridyl, 3H-imidazolyl, indolyl, isoindolyl, indolinyl, isoindolinyl, 1H-indazolyl, 2,3-dihydro-1H-indazolyl, 1H-benzimidazolyl, 2,3-dihydro-1H-benzimidazolyl or 1H-benzotriazolyl group, substituted by an R³ group and optionally substituted by at least one independently selected R⁴ group;

- R³ is selected from a group comprising benzyl, halo-, dihalo- and trihalobenzyl, benzoyl, pyridylmethyl, pyridylmethoxy, phenoxy, benzyloxy, halo-, dihalo- and trihalobenzyloxy and benzenesulphonyl;

or R³ represents trihalomethylbenzyl or trihalomethylbenzyloxy;

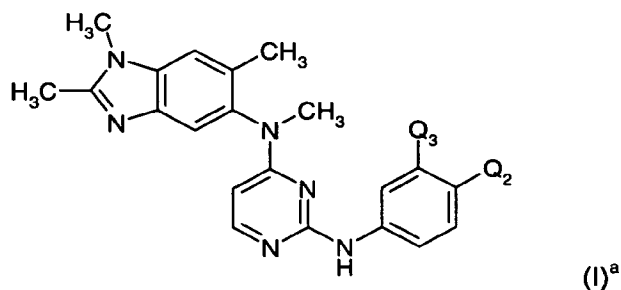
or R³ represents a group of formula



wherein each R⁵ is independently selected from halogen, C₁₋₄ alkyl and C₁₋₄ alkoxy; and n is 0 to 3; and

- each R⁴ is independently hydroxy, halogen, C₁₋₄ alkyl, C₂₋₄ alkenyl, C₂₋₄ alkynyl, C₁₋₄ alkoxy, amino, C₁₋₄ alkylamino, di[C₁₋₄ alkyl]amino, C₁₋₄ alkylthio, C₁₋₄ alkylsulphinyl, C₁₋₄ alkylsulphonyl, C₁₋₄ alkylcarbonyl, carboxy, carbamoyl, C₁₋₄ alkoxycarbonyl, C₁₋₄ alkanoylamino, N-(C₁₋₄ alkyl)carbamoyl, N,N-di(C₁₋₄ alkyl)carbamoyl, cyano, nitro and trifluoromethyl.

2. The method of claim 1, wherein (a) the compound of formula I is a compound of formula I^a:



- 5 or a salt, solvate or physiologically functional derivative thereof;
wherein Q₃ is A¹ when Q₂ is A² and Q₃ is A² when Q₂ is A¹;
wherein

A¹ is hydrogen, halogen, C₁₋₃ alkyl, and

A² is the group defined by $-(Z)_m-(Z^1)-(Z^2)$, wherein

10 Z is CH₂ and m is 0, 1, 2, or 3;

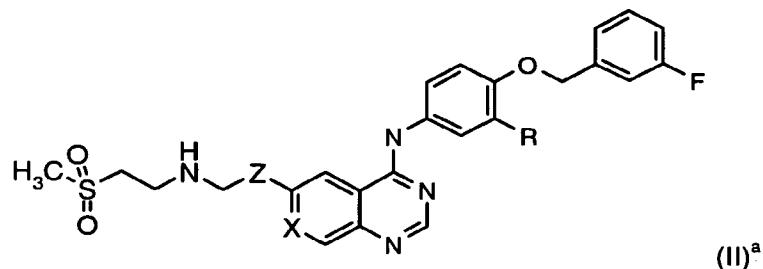
Z¹ is S(O)₂, S(O), or C(O); and

Z² is C₁₋₄ alkyl, NR¹R²

R¹ and R² are each independently selected from hydrogen, or C₁₋₄ alkyl; and

15

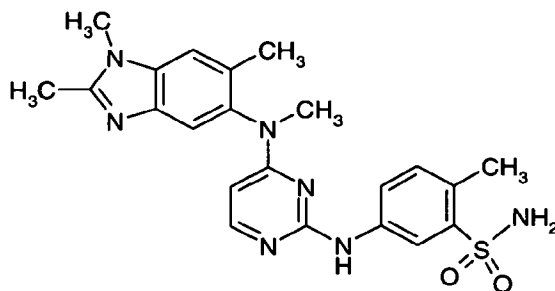
- (b) the compound of formula II is a compound of formula II^a:



- 20 or a salt, solvate or physiologically functional derivative thereof; wherein R is
-Cl or -Br, X is CH, N, or CF, and Z is thiazole or furan.

3. The method of claim 1, wherein (a) the compound of formula I is a compound of formula I^b:

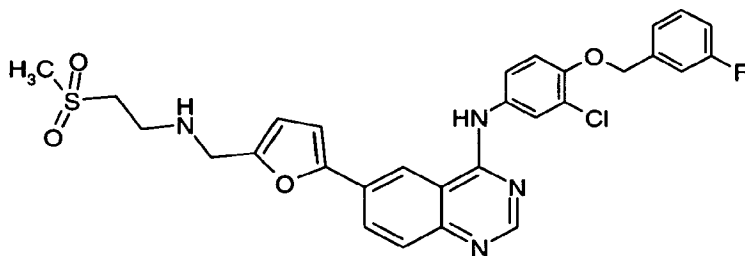
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(I)^b

or a salt, solvate, or physiological functional derivative thereof; and

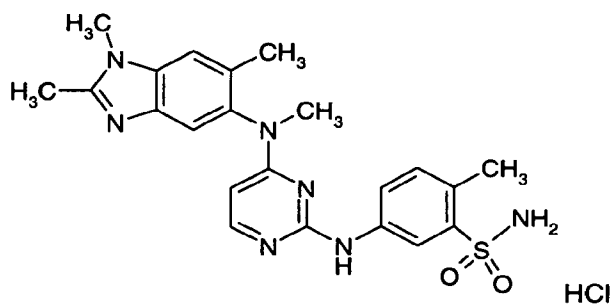
(b) the compound of formula II is a compound of formula II^b:

5

(II)^b

or a salt, solvate, or physiological functional derivative thereof.

- 10 4. The method of claim 1, wherein (a) the compound of formula I is a monohydrochloride salt of a compound of formula I^b:



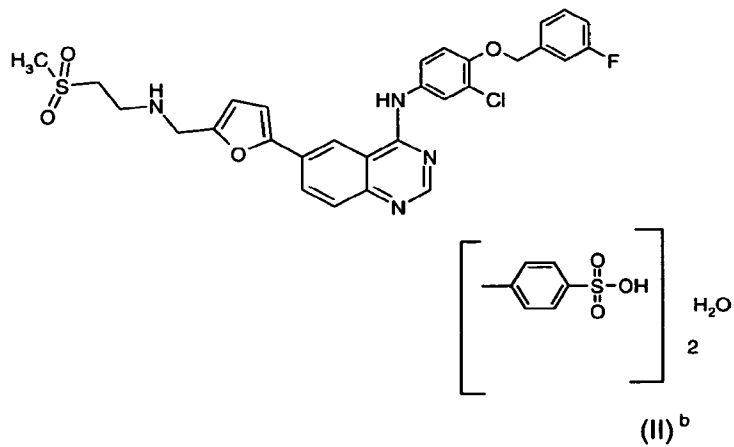
HCl

(I)^b

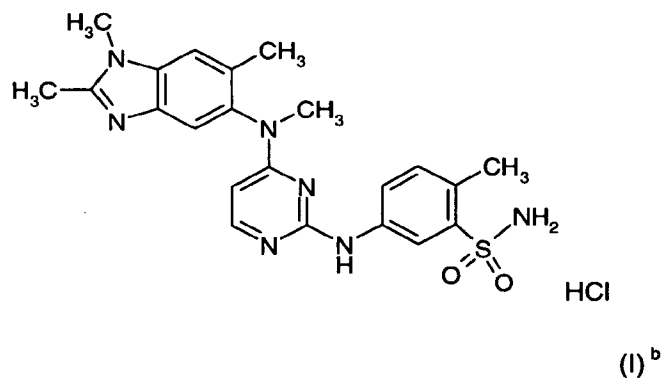
15

; and

(b) the compound of formula II is a monohydrate ditosylate salt of a compound of formula II^b:

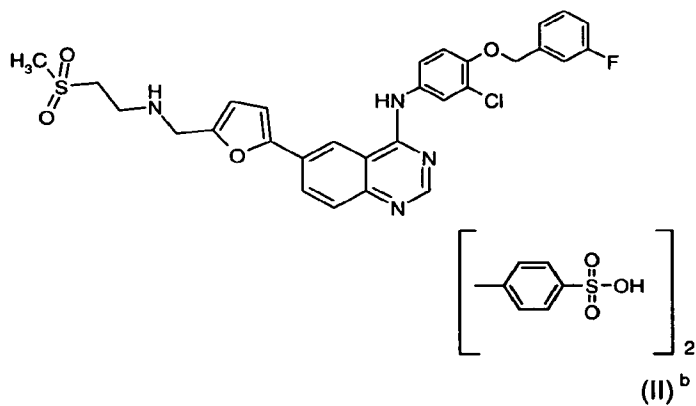


5. The method of claim 1, wherein the compound of formula I is a monohydrochloride salt of a compound of formula I^b:

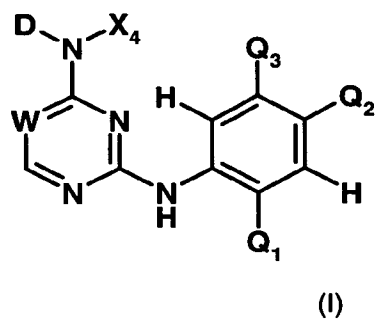


10 ; and

(b) the compound of formula II is an anhydrous ditosylate salt of a compound of formula II^b:

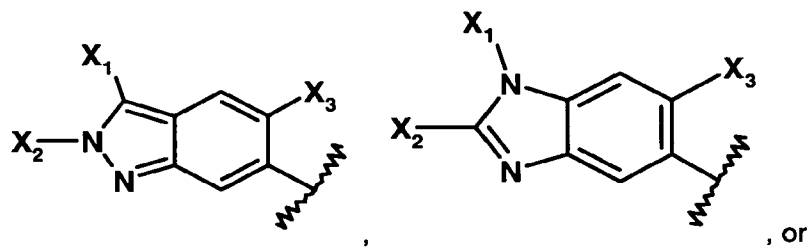


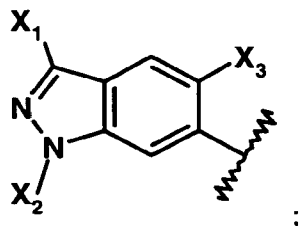
6. A pharmaceutical composition comprising:
 5 (a) a compound of formula I,



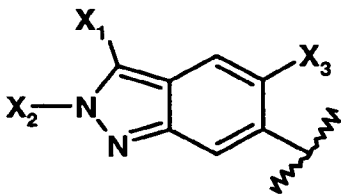
- or a salt, solvate, or physiologically functional derivative thereof;
 10 wherein:

D is



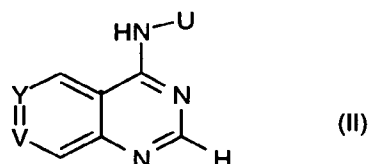


- X_1 is hydrogen, C_{1-4} alkyl, C_{1-4} haloalkyl, or C_{1-4} hydroxyalkyl;
 X_2 is hydrogen, C_{1-4} alkyl, C_{1-4} haloalkyl, $C(O)R^1$, or aralkyl;
 5 X_3 is hydrogen or halogen;
 X_4 is hydrogen, C_{1-4} alkyl, C_{1-4} haloalkyl, heteroaralkyl, cyanoalkyl, $-(CH_2)_pC=CH(CH_2)_tH$, $-(CH_2)_pC\equiv C(CH_2)_tH$, or C_{3-7} cycloalkyl;
 p is 1, 2, or 3;
 t is 0 or 1;
 10 W is N or C-R, wherein R is hydrogen, halogen, or cyano;
 Q_1 is hydrogen, halogen, C_{1-2} haloalkyl, C_{1-2} alkyl, C_{1-2} alkoxy, or C_{1-2} haloalkoxy;
 Q_2 is A^1 or A^2 ;
 Q_3 is A^1 when Q_2 is A^2 and Q_3 is A^2 when Q_2 is A^1 ;
 wherein
 15 A^1 is hydrogen, halogen, C_{1-3} alkyl, C_{1-3} haloalkyl, $-OR^1$, and
 A^2 is the group defined by $-(Z)_m-(Z^1)-(Z^2)$, wherein
 Z is CH_2 and m is 0, 1, 2, or 3, or
 Z is NR^2 and m is 0 or 1, or
 Z is oxygen and m is 0 or 1, or
 20 Z is CH_2NR^2 and m is 0 or 1;
 Z^1 is $S(O)_2$, $S(O)$, or $C(O)$; and
 Z^2 is C_{1-4} alkyl, NR^3R^4 , aryl, arylamino, aralkyl, aralkoxy, or heteroaryl,
 R^1 is C_{1-4} alkyl;
 R^2 , R^3 , and R^4 are each independently selected from hydrogen, C_{1-4} alkyl, C_{3-7}
 25 cycloalkyl, $-S(O)_2R^5$, and $-C(O)R^5$;
 R^5 is C_{1-4} alkyl, or C_{3-7} cycloalkyl; and
 when Z is oxygen then Z^1 is $S(O)_2$ and when D is



then X_2 is C_{1-4} alkyl, C_{1-4} haloalkyl, $C(O)R^1$, or aralkyl; and

(b) a compound of formula II:



- 5 or a salt, solvate, or physiologically functional derivative thereof;
wherein

Y is CR^1 and V is N ;

or Y is CR^1 and V is CR^2 ;

- 10 R^1 represents a group $CH_3SO_2CH_2CH_2NHCH_2-Ar-$, wherein Ar is selected from phenyl, furan, thiophene, pyrrole and thiazole, each of which may optionally be substituted by one or two halo, C_{1-4} alkyl or C_{1-4} alkoxy groups;

R^2 is selected from the group comprising hydrogen, halo, hydroxy, C_{1-4} alkyl, C_{1-4}

- 15 alkoxy, C_{1-4} alkylamino and $di[C_{1-4}$ alkyl]amino;

U represents a phenyl, pyridyl, 3H-imidazolyl, indolyl, isoindolyl, indolinyl, isoindolinyl, 1H-indazolyl, 2,3-dihydro-1H-indazolyl, 1H-benzimidazolyl, 2,3-dihydro-1H-benzimidazolyl or 1H-benzotriazolyl group, substituted by an R^3 group and optionally substituted by at least one independently selected R^4 group;

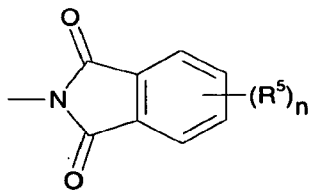
- 20

R^3 is selected from a group comprising benzyl, halo-, dihalo- and trihalobenzyl, benzoyl, pyridylmethyl, pyridylmethoxy, phenoxy, benzyloxy, halo-, dihalo- and trihalobenzyloxy and benzenesulphonyl;

- 25

or R^3 represents trihalomethylbenzyl or trihalomethylbenzyloxy;

or R^3 represents a group of formula

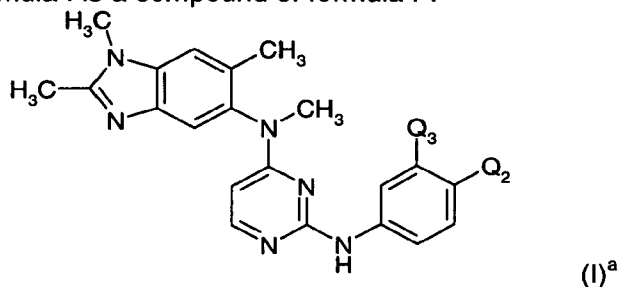


wherein each R^5 is independently selected from halogen, C_{1-4} alkyl and C_{1-4} alkoxy; and n is 0 to 3; and

- 5 each R^4 is independently hydroxy, halogen, C_{1-4} alkyl, C_{2-4} alkenyl, C_{2-4} alkynyl, C_{1-4} alkoxy, amino, C_{1-4} alkylamino, di[C_{1-4} alkyl]amino, C_{1-4} alkylthio, C_{1-4} alkylsulphinyl, C_{1-4} alkylsulphonyl, C_{1-4} alkylcarbonyl, carboxy, carbamoyl, C_{1-4} alkoxycarbonyl, C_{1-4} alkanoylamino, N -(C_{1-4} alkyl)carbamoyl, N,N -di(C_{1-4} alkyl)carbamoyl, cyano, nitro and trifluoromethyl.

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7. The pharmaceutical composition of claim 6, wherein (a) the compound of formula I is a compound of formula I^a:



15

or a salt, solvate or physiologically functional derivative thereof; wherein Q_3 is A^1 when Q_2 is A^2 and Q_3 is A^2 when Q_2 is A^1 ; wherein

A^1 is hydrogen, halogen, C_{1-3} alkyl, and

A^2 is the group defined by $-(Z)_m-(Z^1)-(Z^2)$, wherein

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Z is CH_2 and m is 0, 1, 2, or 3;

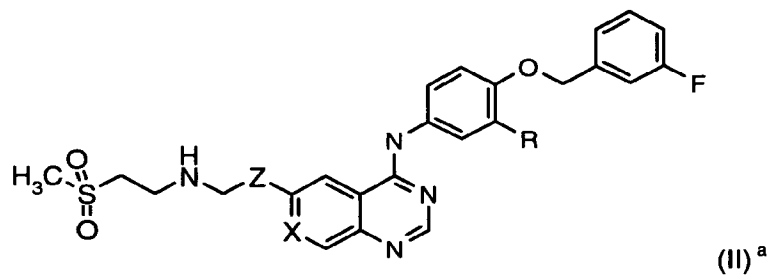
Z^1 is $S(O)_2$, $S(O)$, or $C(O)$; and

Z^2 is C_{1-4} alkyl, NR^1R^2

R^1 and R^2 are each independently selected from hydrogen, or C_{1-4} alkyl; and

25

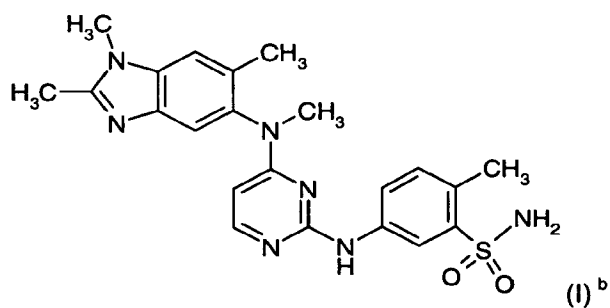
(b) the compound of formula II is a compound of formula II^a:



or a salt, solvate or physiologically functional derivative thereof; wherein R is -Cl or -Br, X is CH, N, or CF, and Z is thiazole or furan.

5

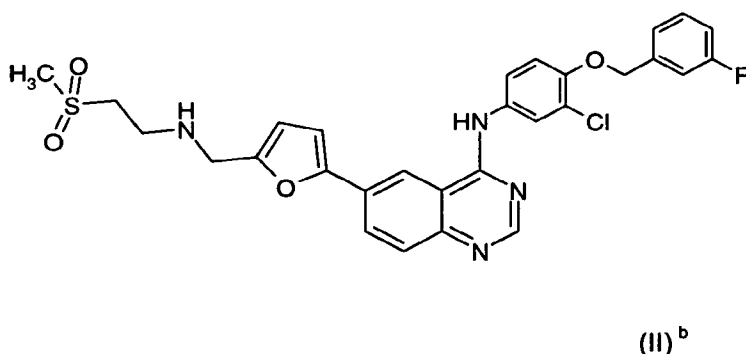
8. The pharmaceutical composition of claim 6, wherein (a) the compound of formula I is a compound of formula I^b:



or a salt, solvate, or physiological functional derivative thereof; and

10

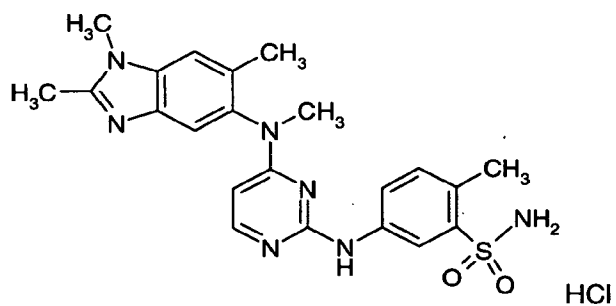
- (b) the compound of formula II is a compound of formula II^b



or a salt, solvate, or physiological functional derivative thereof.

15

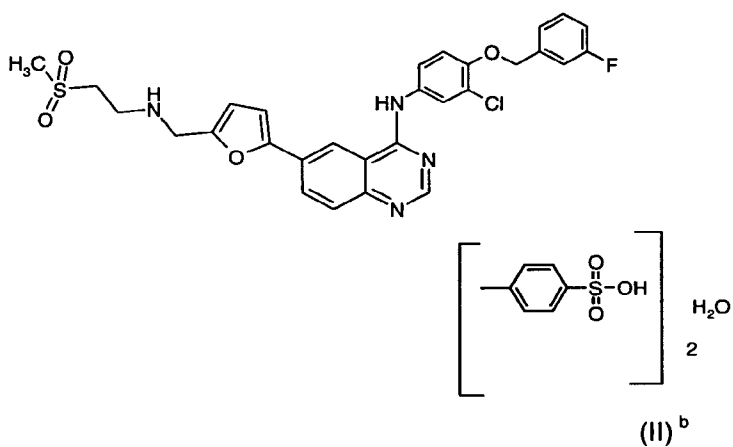
9. The pharmaceutical composition of claim 6, wherein (a) the compound of formula I is a monohydrochloride salt of a compound of formula I^b:

(I)^b

; and

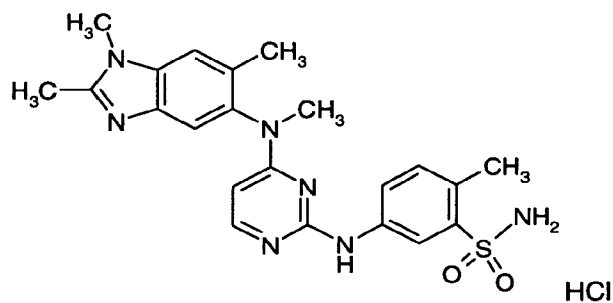
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(b) the compound of formula II is a monohydrate ditosylate salt of the compound of formula II^b:



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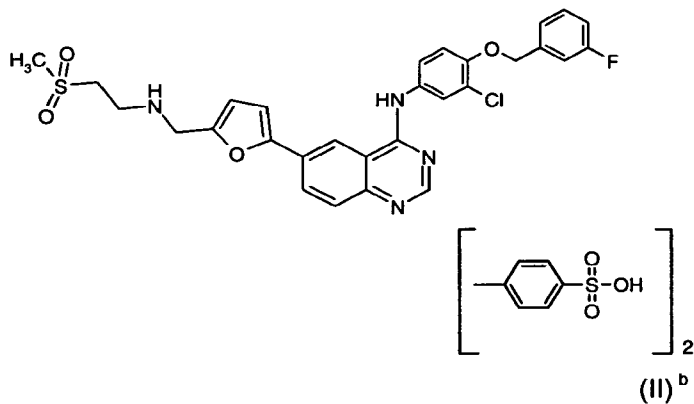
10. The pharmaceutical composition of claim 6, wherein (a) the compound of formula I is a monohydrochloride salt of a compound of formula I^b:

(I)^b

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; and

(b) the compound of formula II is an anhydrous ditosylate salt of the compound of formula II^b:



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11. A pharmaceutical combination comprising: a compound of formula I, I^a or I^b or salt, solvate or physiologically functional derivative thereof, and a compound of formula II, II^a or II^b or salt, solvate or physiologically functional derivative thereof for use in therapy.
12. The use of a pharmaceutical combination comprising: a compound of formula I, I^a or I^b or salt, solvate or physiologically functional derivative thereof, and a compound of formula II, II^a or II^b or salt, solvate or physiologically functional derivative thereof for the preparation of a medicament useful in the treatment of cancer.

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ABSTRACT

A method of treating cancer is described including administration of a pyrimidine derivative and a quinazoline derivative as well as a pharmaceutical composition including the same.

Figure 1

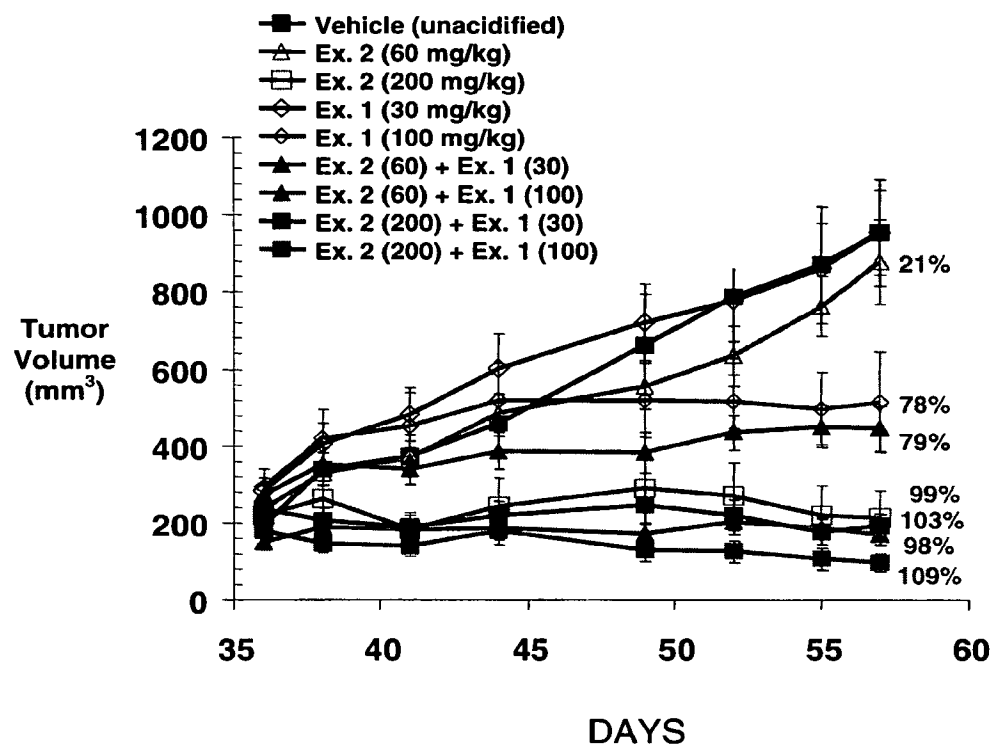


Table 1

Percent Inhibition of BT474 Tumor growth by Example 1 and Example 2.

	Exp. 1	Exp. 2
Example 2 (60 mg/kg)	21	58*
Example 2 (200 mg/kg)	99*	85*
Example 1 (30 mg/kg)	16	47*
Example 1 (100 mg/kg)	78*	64*
Ex. 1 (30 mg/kg) + Ex. 2 (60 mg/kg)	79*	71*
Ex. 1 (100 mg/kg) + Ex. 2 (60 mg/kg)	98*	84*
Ex. 1 (30 mg/kg) + Ex. 2 (200 mg/kg)	103*	91*
Ex. 1 (100 mg/kg) + Ex. 2 (200 mg/kg)	109*	88*

* Significant difference from vehicle treated group (paired comparison of slopes using Tukey-Kramer HSD)

Table 2

Percent Inhibition of NCI H322 Tumor growth by Example 1 and Example 2.

	Exp. 1	Exp. 2	Exp. 3
Example 2 (20 mg/kg)	N.D	31	28
Example 2 (60 mg/kg)	75*	N.D.	54
Example 1 (30 mg/kg)	34	20	86*
Example 1 (100 mg/kg)	56*	62	88*
Ex. 1 (30 mg/kg) + Ex. 2 (20 mg/kg)	N.D.	43	60*
Ex. 1 (100 mg/kg) + Ex. 2 (20 mg/kg)	N.D.	69*	94*
Ex. 1 (30 mg/kg) + Ex. 2 (60 mg/kg)	81*	N.D.	92*
Ex. 1 (100 mg/kg) + Ex. 2 (60 mg/kg)	81*	N.D.	108*

N.D. - Experiment not done

* Significant difference from vehicle treated group (paired comparison of slopes using Tukey-Kramer HSD)